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Carbazole-based small molecule electron donors: syntheses, characterization, and material properties

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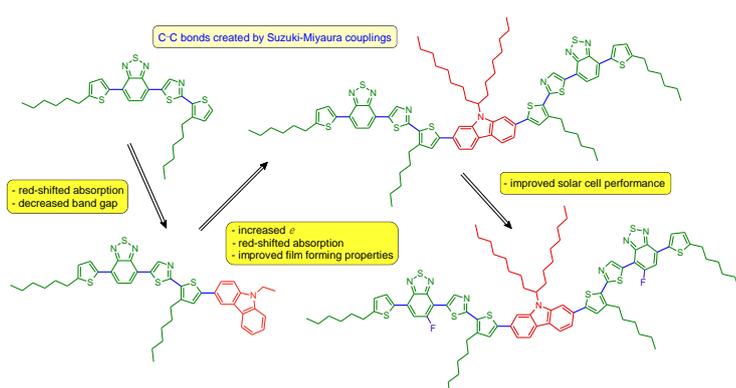
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Graphical abstract



Keywords

Absorption, DFT, Electron donor, Organic solar cell, Suzuki-Miyaura, Synthesis

Highlights

- Novel *D-A* and *A-D-A* type small molecule electron donors were synthesized
- Characterization by computational, electrochemical, and spectroscopic methods
- *A-D-A* type materials were tested as active layer components in OSC devices
- Relations between molecular structures and properties are studied and discussed

Abstract

Efficient synthetic methods for carbazole-based small molecule electron donors with donor–acceptor ($D-A$) and $A-D-A$ type structures were developed. In order to study the relation between chemical structures and material properties, the prepared compounds were characterized in detail using absorption spectroscopy, differential pulse voltammetry, and computational methods. In addition, symmetrical $A-D-A$ type compounds were tested as an active layer component in bulk heterojunction based organic solar cell (OSC) devices with conventional structure. The results show that the two compound types have many similar properties. However, the extended molecular structure of $A-D-A$ type compounds offer better film forming properties and higher molar absorption coefficients compared with the $D-A$ type materials. Furthermore, the attachment of fluoro substituents in the A units has a positive effect on all solar cell device parameters. Moreover, the computational studies revealed that the molecular structures are twisted between the central carbazole D unit and π -bridge which may result in inefficient intramolecular charge transfer and, also, relatively limited short-circuit currents in OSC devices.

1. Introduction

In the recent years, solution processed small molecule donors have attracted attention as alternatives to conjugated polymer-based donors in organic solar cells (OSCs). These materials show several advantages compared to their polymeric counterparts, such as well-defined structures without the end group contaminants, simple synthesis and purification, low dispersity, and excellent batch-to-batch reproducibility [1,2,3,4,5,6,7].

One of the most efficient structures of small molecule donor materials has turned out to be the $A-D-A$ type structure in which A refers to an electron acceptor unit and D refers to an electron donor unit. Often, these kinds of molecules are also termed *push-pull-push* type donor materials. By changing both the central D and terminal A units and a possible π -conjugation bridge in between the

A and *D* units, desired material properties can be tailored and high power conversion efficiencies (PCEs) achieved [8]. For example, one of the best-performing small molecule donors is p-DTS(FBTTh₂)₂ (Fig. 1) [5]. Up to 8.9% PCE has been reported with this molecule by optimizing the OSC device structure [9]. Moreover, it has been demonstrated that some small molecule materials can produce very high PCEs both in bulk heterojunction (BHJ) based OSCs and in perovskite solar cells [10].

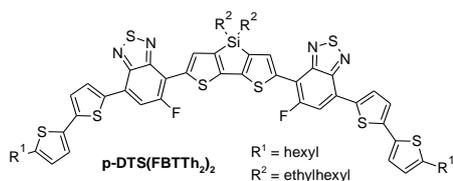


Fig. 1. The chemical structure of p-DTS(FBTTh₂)₂.

Especially, 2,1,3-benzothiadiazole (BT) unit and its substituted derivatives have been widely utilized as building blocks (*A* units) in both polymer and small molecule electron donor materials [11,12]. Moreover, various *D* units have been studied as a component of OSC donor materials [12,13]. Carbazole (Cz) has been employed as the *D* unit in various semiconducting polymers and dye molecules [14,15]. The main advantage of the poly(2,7-carbazole)-based OSC devices is high open circuit voltages (V_{oc}). Moreover, internal quantum efficiency can be close to 100% [16]. Few years ago, Cz-based small molecule materials showed promising results as donor components in BHJ OSCs [17,18]. Since then, small molecule materials containing Cz as the *D* unit have gained growing interest and shown promising results. At least, three main structures *A–D–A* [19,20], *D–A* [21,22], and *D–A–D* [23,24,25,26] have been introduced. However, more experimental studies are needed to discover new state-of-the-art small molecule electron donors and, in particular, to gain systematic information about the relationships between their chemical structures, material properties, and performance in OSC applications.

Recently, we reported a synthetic pathway to BT-based building blocks [27]. Since then, we have further developed the synthetic strategy and, here, we present the syntheses of two

unsymmetrical *D*–*A* and two symmetrical *A*–*D*–*A* type small molecules. In these compounds, Cz acts as a *D* unit and either BT or its fluoro-substituted derivative as an *A* unit. Thiophene and thiazole have been used as a π -bridge in between the *D* and *A* units. In order to find out how the changes in molecular structures affect the material properties, the prepared *D*–*A* and *A*–*D*–*A* type small molecule compounds have been characterized by spectroscopic, electrochemical, and computational methods. Moreover, the symmetrical *A*–*D*–*A* type compounds have been tested as electron donor materials in BHJ-based OSC devices.

2. Experimental section

Commercial reagents were used as received. 2-(3-Hexylthiophen-2-yl)-5-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)thiazole and compound **3b** were synthesized using the previously published methods [27]. The chemical structures of new compounds were characterized by using NMR and HRMS techniques. Melting points (reported as peak values) of compounds **5a**, **5b**, **6a**, and **6b** were measured by using Mettler Toledo DSC 1 apparatus with a TSO800GC1 Gas Control system. Samples of 1.6–2.2 mg were placed in 40 μ L Al crucibles with pierced lids. The DSC scans were carried out from 5 $^{\circ}$ C to 300 $^{\circ}$ C at the heating rate of 20 $^{\circ}$ C min^{-1} under a nitrogen flow of 60 mL min^{-1} .

2.1. Syntheses

2.1.1. Synthesis of 4-bromo-5-fluoro-7-(5-hexylthiophen-2-yl)-2,1,3-benzothiadiazole (**2**)

Toluene (6 mL), DMA (6 mL), distilled water (1.5 mL), and 5-hexyl-2-thiopheneboronic acid pinacol ester (1.02 equiv, 140.6 mg, 0.48 mmol) were deoxygenated with argon for 15 min in a reaction tube with a magnetic stirring bar. 4,7-Dibromo-5-fluoro-2,1,3-benzothiadiazole (**1**) (146.9 mg, 0.47 mmol), Cs_2CO_3 (2.5 equiv, 384.1 mg, 1.18 mmol), $\text{Pd}(\text{OAc})_2$ (2.6 mol%, 2.8 mg, 12 μ mol), and Xantphos (2.6 mol%, 6.8 mg, 12 μ mol) were added. The sealed tube was evacuated and backfilled with argon five times. The reaction mixture was stirred and heated in an oil bath (60 $^{\circ}$ C) for 3 h. The

reaction mixture was filtered through a thin pad of silica gel rinsing with toluene and evaporated under reduced pressure. The product was purified by using flash chromatography (SiO₂, toluene 1 : 1 *n*-heptane). The isolated product **2** was collected as a yellow solid (186.7 mg) in >99% yield. Mp 90 °C. ¹H NMR (400 MHz, CDCl₃) δ ppm 0.91 (t, *J*=6.9 Hz, 3H), 1.32–1.38 (m, 4H), 1.38–1.46 (m, 2H), 1.75 (quin, *J*=7.5 Hz, 2H), 2.89 (t, *J*=7.7 Hz, 2H), 6.88 (d, *J*=3.7 Hz, 1H), 7.62 (d, *J*=10.0 Hz, 1H), 7.95 (d, *J*=3.7 Hz, 1H). HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₁₆H₁₇N₂S₂BrF 399.0001; Found 398.9999.

2.1.2. Synthesis of compound **3a**

Toluene (2.5 mL), DMA (2.5 mL), and distilled water (0.5 mL) were deoxygenated with argon for 15 min in a reaction tube with a magnetic stirring bar. Compound **2** (99.9 mg, 0.25 mmol), 2-(3-hexylthiophen-2-yl)-5-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)thiazole (1.04 equiv, 99.2 mg, 0.26 mmol), Cs₂CO₃ (2.5 equiv, 204.9 mg, 0.63 mmol), *t*-Bu₃P·HBF₄ (12 mol%, 8.7 mg, 30 μmol) and Pd₂(dba)₃ (3 mol%, 6.9 mg, 7.5 μmol) were added. The sealed tube was evacuated and backfilled with argon five times. The reaction mixture was stirred and heated in an oil bath (90 °C) for 21 h. The reaction mixture was filtered through a thin pad of silica gel rinsing with toluene and evaporated under reduced pressure. The product was purified by using flash chromatography (SiO₂, toluene). The isolated product **3a** was collected as a red solid (100.4 mg) in 70% yield. Mp 79 °C. ¹H NMR (400 MHz, CDCl₃) δ ppm 0.90–0.94 (m, 6H), 1.33–1.54 (m, 12H), 1.73–1.81 (m, 4H), 2.90 (t, *J*=7.6 Hz, 2H), 3.05 (t, *J*=7.9 Hz, 2H), 6.90 (d, *J*=3.7 Hz, 1H), 7.02 (d, *J*=5.1 Hz, 1H), 7.35 (d, *J*=5.1 Hz, 1H), 7.70 (d, *J*=13.0 Hz, 1H), 8.00 (d, *J*=3.7 Hz, 1H), 8.90 (s, 1H). HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₃N₃S₄F 570.1541; Found 570.1534.

2.1.3. Synthesis of compound **4a**

Compound **3a** (80.2 mg, 0.14 mmol) was dissolved in CH₂Cl₂ (10 mL) and NBS (1.1 equiv, 27.8 mg, 0.16 mmol) was added. The reaction mixture was stirred at room temperature for 20 h. The

solvent was evaporated and the crude product was subjected to flash chromatography (SiO₂, toluene). The pure compound **4a** was isolated as a red solid (80.0 mg) in 88% yield. Mp 96 °C. ¹H NMR (400 MHz, CDCl₃) δ ppm 0.90–0.95 (m, 6H), 1.32–1.51 (m, 12H), 1.69–1.80 (m, 4H), 2.90 (t, *J*=7.6 Hz, 2H), 2.95 (t, *J*=7.8 Hz, 2H), 6.89 (d, *J*=3.7 Hz, 1H), 6.96 (s, 1H), 7.67 (d, *J*=13.0 Hz, 1H), 7.98 (d, *J*=3.7 Hz, 1H), 8.84 (s, 1H). HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₂N₃S₄BrF 648.0646; Found 648.0643.

2.1.4. Synthesis of compound **4b**

The synthesis was carried out using the same procedure as for compound **4a**. The specific amounts of reagents used were: compound **3b** (71.7 mg, 0.130 mmol), NBS (1.02 equiv, 23.5 mg, 0.132 mmol), and CH₂Cl₂ (4.8 mL). Pure compound **4b** was isolated as a red solid (42.9 mg) in 52% yield. Mp 130 °C. ¹H NMR (400 MHz, CDCl₃) δ ppm 0.90–0.95 (m, 6H), 1.32–1.51 (m, 12H), 1.70–1.80 (m, 4H), 2.90 (t, *J*=7.7 Hz, 2H), 2.96 (t, *J*=7.8 Hz, 2H), 6.90 (d, *J*=3.8 Hz, 1H), 6.97 (s, 1H), 7.82 (d, *J*=7.6 Hz, 1H), 7.85 (d, *J*=7.6 Hz, 1H), 7.99 (d, *J*=3.8 Hz, 1H), 8.69 (s, 1H). ¹³C NMR (100.6 MHz, CDCl₃) δ ppm 14.1, 14.1, 22.6, 22.6, 28.8, 29.2, 29.8, 30.2, 30.3, 31.6, 31.6, 114.8, 122.3, 124.8, 125.4, 126.4, 127.2, 128.1, 133.2, 133.5, 134.1, 136.4, 141.7, 142.8, 148.6, 152.3, 152.5, 159.9. HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₃N₃S₄Br 630.0741; Found 630.0746. A small amount of byproduct **7** was isolated for analyses. Mp 90 °C. ¹H NMR (400 MHz, CDCl₃) δ ppm 0.90–0.94 (m, 6H), 1.32–1.51 (m, 12H), 1.70–1.79 (m, 4H), 2.87 (t, *J*=7.5 Hz, 2H), 2.96 (t, *J*=7.8 Hz, 2H), 6.88 (s, 1H), 6.98 (s, 1H), 7.93 (d, *J*=7.6 Hz, 1H), 8.07 (d, *J*=7.6 Hz, 1H), 8.74 (s, 1H). ¹³C NMR (100.6 MHz, CDCl₃) δ ppm 14.1, 14.1, 22.6, 22.6, 28.8, 29.2, 29.7, 30.2, 30.2, 31.1, 31.5, 31.6, 109.5, 115.0, 124.2, 125.5, 125.8, 128.3, 129.8, 129.9, 133.3, 133.5, 133.7, 142.3, 143.0, 148.2, 152.0, 153.5, 160.4. HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₂₉H₃₂N₃S₄Br₂ 707.9846; Found 707.9836.

2.1.5. Synthesis of compound **5a**

Toluene (2.25 mL), DMA (2.25 mL), and distilled water (0.5 mL) were deoxygenated with argon for 15 min in a reaction tube with a magnetic stirring bar. Compound **4a** (75.9 mg, 0.117 mmol), 9-ethylcarbazole-3-boronic acid (1.05 equiv, 29.3 mg, 0.123 mmol), Cs₂CO₃ (2.5 equiv, 96.1 mg, 0.29 mmol), Pd(OAc)₂ (6 mol%, 1.5 mg, 6.7 μmol), and Xantphos (5 mol%, 3.4 mg, 5.9 μmol) were added. The sealed tube was evacuated and backfilled with argon five times. The reaction mixture was stirred and heated in an oil bath (100 °C) for 1.5 h. The reaction mixture was filtered through a thin pad of silica gel rinsing with toluene and evaporated under reduced pressure. The product was purified by using flash chromatography (SiO₂, toluene). The solid product was boiled in acetone-methanol mixture (1:2), cooled to room temperature, filtered and washed with acetone-methanol mixture several times. The isolated product **5a** was collected as a deep dark red solid (87.2 mg) in 98% yield. Mp 102 °C. ¹H NMR (400 MHz, CD₂Cl₂) δ ppm 0.91–0.98 (m, 6H), 1.33–1.49 (m, 13H), 1.57 (quin, *J*=7.1 Hz, 2H), 1.75 (quin, *J*=7.5 Hz, 2H), 1.84 (quin, *J*=7.6 Hz, 2H), 2.88 (t, *J*=7.6 Hz, 2H), 3.04 (t, *J*=7.8 Hz, 2H), 4.35 (q, *J*=7.2 Hz, 2H), 6.87 (d, *J*=3.7 Hz, 1H), 7.24–7.28 (m, 2H), 7.39–7.43 (m, 2H), 7.47–7.51 (m, 1H), 7.65 (d, *J*=13.1 Hz, 1H), 7.75 (dd, *J*=8.6, 1.8 Hz, 1H), 7.95 (d, *J*=3.7 Hz, 1H), 8.12 (d, *J*=7.6 Hz, 1H), 8.34 (d, *J*=1.4 Hz, 1H), 8.84 (s, 1H). HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₄₃H₄₄N₄S₄F 763.2433; Found 763.2441.

2.1.6. Synthesis of compound **5b**

The synthesis was carried out using the same procedure as for compound **5a**. The specific amounts of reagents used were: toluene (1.5 mL), DMA (1.5 mL), distilled water (0.38 mL), compound **4b** (50.2 mg, 0.080 mmol), 9-ethylcarbazole-3-boronic acid (1.04 equiv, 19.9 mg, 0.083 mmol), Cs₂CO₃ (2.5 equiv, 64.8 mg, 0.20 mmol), Pd(OAc)₂ (8 mol%, 1.5 mg, 6.7 μmol), and Xantphos (5 mol%, 2.5 mg, 4.3 μmol). Pure compound **5b** was isolated as a dark red solid (57.1 mg) in 96% yield. Mp 178 °C. ¹H NMR (400 MHz, CD₂Cl₂) δ ppm 0.91–0.97 (m, 6H), 1.34–1.48 (m, 13H), 1.58 (quin, *J*=7.3 Hz 2H), 1.77 (quin, *J*=7.5 Hz, 2H), 1.86 (quin, *J*=7.6 Hz, 2H), 2.91 (t, *J*=7.6 Hz, 2H), 3.08 (t, *J*=8.0 Hz, 2H), 4.40 (q, *J*=7.2 Hz, 2H), 6.91 (d, *J*=3.7 Hz, 1H), 7.25–7.32 (m, 2H),

7.45–7.53 (m, 3H), 7.80 (dd, $J=8.6, 1.8$ Hz, 1H), 7.83 (d, $J=7.6$ Hz, 1H), 7.87 (d, $J=7.6$ Hz, 1H), 7.99 (d, $J=3.7$ Hz, 1H), 8.16 (d, $J=7.7$ Hz, 1H), 8.41 (d, $J=1.6$ Hz, 1H), 8.75 (s, 1H). ^{13}C NMR (100.6 MHz, CDCl_3) δ ppm 13.8, 14.1, 14.1, 22.6, 22.7, 28.9, 29.5, 29.9, 30.3, 30.6, 31.6, 31.6, 31.8, 37.7, 108.7, 108.8, 117.7, 119.3, 120.6, 122.8, 122.9, 123.5, 123.9, 124.8, 124.9, 125.2, 125.3, 126.1, 126.7, 127.9, 129.9, 133.3, 136.6, 139.9, 140.5, 142.0, 143.9, 146.6, 148.3, 152.4, 152.5, 161.3. HRMS (ESI+, TOF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{43}\text{H}_{45}\text{N}_4\text{S}_4$ 745.2527; Found 745.2524.

2.1.7. Synthesis of compound **6a**

Toluene (3 mL), DMA (3 mL), and distilled water (0.6 mL) were deoxygenated with argon for 15 min in a reaction tube with a magnetic stirring bar. Compound **4a** (2.05 equiv, 98.7 mg, 0.152 mmol), 9-(9-heptadecanyl)-2,7-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)carbazole (48.8 mg, 0.074 mmol), Cs_2CO_3 (5 equiv, 124.0 mg, 0.381 mmol), $\text{Pd}(\text{OAc})_2$ (10 mol%, 1.7 mg, 7.6 μmol), and Xantphos (10 mol%, 4.4 mg, 7.6 μmol) were added. The sealed tube was evacuated and backfilled with argon five times. The reaction mixture was stirred and heated in an oil bath (100 $^\circ\text{C}$) for 4 h. The reaction mixture was filtered through a thin pad of silica gel rinsing with CHCl_3 and evaporated under reduced pressure. The product was purified by using flash chromatography (SiO_2 , CHCl_3). The solid product was boiled in acetone (3 mL), filtered and washed with hot acetone (9 mL). The procedure was repeated four times. The isolated product **6a** was collected as a deep dark solid (109.3 mg) in 96% yield. Mp 156 $^\circ\text{C}$. ^1H NMR (400 MHz, CDCl_3 , at 330 K) δ ppm 0.80–0.84 (t, $J=6.6$, 6H), 0.92–0.99 (m, 12H), 1.14–1.50 (m, 44H) [28], 1.60 (quin, $J=7.3$ Hz, 4H), 1.79 (quin, $J=7.3$ Hz, 4H), 1.89 (quin, $J=7.6$ Hz, 4H), 2.01–2.10 (m, 2H), 2.34–2.44 (m, 2H), 2.92 (t, $J=7.5$ Hz, 4H), 3.12 (t, $J=7.6$ Hz, 4H), 4.66 (tt, $J=9.7, 4.9$ Hz, 1H), 6.91 (d, $J=3.7$ Hz, 2H), 7.33 (s, 2H), 7.56 (d, $J=8.1$ Hz, 2H), 7.72 (d, $J=12.8$ Hz, 2H), 7.80 (br s, 2H), 8.02 (d, $J=3.7$ Hz, 2H), 8.07 (d, $J=8.1$ Hz, 2H), 8.94 (s, 2H). HRMS (ESI+, TOF) m/z : $[\text{M}+\text{H}]^+$ Calcd for $\text{C}_{87}\text{H}_{104}\text{N}_7\text{F}_2\text{S}_8$ 1540.6087; Found 1540.6097.

2.1.8. Synthesis of compound **6b**

The synthesis was carried out using the same procedure as for compound **6a**. The specific amounts of reagents used were: toluene (2.2 mL), DMA (2.2 mL), distilled water (0.4 mL), compound **4b** (2 equiv, 70.2 mg, 0.11 mmol), 9-(9-heptadecanyl)-2,7-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)carbazole (35.9 mg, 0.055 mmol), Cs₂CO₃ (5 equiv, 90.2 mg, 0.277 mmol), Pd(OAc)₂ (11 mol%, 1.4 mg, 6.2 μmol), and Xantphos (10 mol%, 3.2 mg, 5.6 μmol). The isolated product **6b** was collected as a deep dark solid (64.3 mg) in 78% yield. Mp 159 °C. ¹H NMR (400 MHz, CDCl₃, at 325 K) δ ppm 0.79 (t, *J*=6.9 Hz, 6H), 0.89–0.96 (m, 12H), 1.17–1.23 (m, 18H), 1.29–1.36 (m, 14H), 1.39–1.47 (m, 12H), 1.56 (quin, *J*=7.2 Hz, 4H), 1.75 (quin, *J*=7.5 Hz, 4H), 1.84 (quin, *J*=7.6 Hz, 4H), 1.99–2.07 (m, 2H), 2.32–2.41 (m, 2H), 2.87 (t, *J*=7.6 Hz, 4H), 3.05 (t, *J*=7.7 Hz, 4H), 4.63 (tt, *J*=9.7, 5.0 Hz, 1H), 6.85 (d, *J*=3.7 Hz, 2H), 7.27 (s, 2H), 7.50 (d, *J*=8.1 Hz, 2H), 7.68–7.80 (m, 6H), 7.95 (d, *J*=3.7 Hz, 2H), 8.01 (d, *J*=8.1 Hz, 2H), 8.73 (s, 2H). ¹³C NMR (100.6 MHz, CDCl₃, at 325 K) δ ppm 14.0, 14.0, 14.1, 22.6, 22.7, 27.0, 28.9, 29.2, 29.4, 29.4, 29.5, 30.0, 30.4, 30.6, 31.6, 31.6, 31.8, 31.8, 34.0, 56.9, 106.3, 108.9, 117.3, 120.7, 120.7, 122.8, 124.8, 125.3, 126.3, 126.4, 127.1, 128.1, 131.0, 131.3, 133.7, 136.6, 142.1, 143.9, 146.5, 148.4, 152.5, 152.6, 161.2. HRMS (ESI+, TOF) *m/z*: [M+H]⁺ Calcd for C₈₇H₁₀₆N₇S₈ 1504.6275; Found 1504.6283.

2.2. Spectroscopic measurements and electrochemical characterization

The steady-state absorption spectra were measured with a Shimadzu UV-3600 UV/Vis/NIR spectrophotometer both in solution and in thin films. Thin films of the three compounds were deposited by spin-coating (WS-400B-6NPP/LITE, Laurell Technologies) from CHCl₃ solution (2000 rpm, 2000 rpm/s, 1 min) onto clean quartz substrates.

Differential pulse voltammetry (DPV), for HOMO/LUMO determination of the target compounds, was performed by employing a potentiostat (Compact-Stat, Ivium Technologies) and a three-electrode cell configuration. Dry tetrabutyl ammonium tetrafluoroborate in dichloromethane (0.1 M) was the supporting electrolyte, glass platinum electrode the working electrode, Pt wire the

counter-electrode, and Ag/AgCl wire the pseudo-reference electrode. Ferrocene/ferrocenium (Fc/Fc⁺) couple was used as an internal standard reference to scale the measured potentials against the vacuum level [29]. All solutions were deoxygenated with N₂ prior to each experiment. The measurements were carried out between -2.4 V and 1.6 V, scanning in both directions with scan rate of 50 mV/s. The HOMO and LUMO energy levels were calculated from the oxidation potentials observed from the DPV curves according to the equations:

$$E_{\text{HOMO}}(\text{eV}) = - (E_{\text{ox}} - E_{\text{Fc/Fc}^+}) - 4.80 \text{ eV} \quad (1)$$

$$E_{\text{LUMO}}(\text{eV}) = (E_{\text{red}} - E_{\text{Fc/Fc}^+}) - 4.80 \text{ eV} \quad (2)$$

where E_{ox} and E_{red} are the oxidation and reduction potentials of the sample and $E_{\text{Fc/Fc}^+}$ the potential of ferrocene. E_{ox} and $E_{\text{Fc/Fc}^+}$ are both referred against the Ag/AgCl reference electrode. The value -4.80 eV is the energy level of ferrocene against vacuum [30].

2.3. Computations

All the calculations were carried out with the Gaussian 09 (Revision D.01) suite of programs [31]. The optimal dihedral angles (see Fig. S4) between adjoining units present in compounds **5a**, **5b**, **6a**, and **6b** were determined with the relaxed potential energy scans (PES) at 5° intervals, i.e. the selected dihedral angle was fixed while optimizing the geometry of the two-unit-entity otherwise fully with the density functional theory (DFT) at the B3LYP [32,33]/6-31+G** level of theory. Only two units of the whole molecules were included per a PES scan due to a computation demand of the diffuse basis set. In the PES studies, the ethyl and hexyl side chains were replaced with methyl groups. Additional conformational studies were also carried out for compounds **5a** and **6a** (see Supplementary material for further information). The dihedral angles of compounds **5a**, **5b**, **6a**, and **6b** were set according to the PES scans and then the ground-state geometries of the whole molecules with the longer ethyl and hexyl side chains (Scheme 2) were optimized both in vacuum and chloroform at the B3LYP/6-31G** level of theory. Solvation effects were taken into account by means of the integral equation formalism of the polarizable continuum model using the dielectric constant of 4.71 for

chloroform [34,35]. The final optimized geometries (Fig. 3) were confirmed at the same level of theory to be the minimum energy structures by the absence of the imaginary frequency. The vertical transition energies and oscillator strengths of the compounds were determined both in vacuum and chloroform with TDDFT for the first 20 excited states at the ω B97XD/6-31G** levels of theory. The solvent TDDFT calculations were carried out using the geometries optimized in chloroform. The electronic transitions were qualitatively described using NTOs as a representation for the transition density matrix [36]. Pictorial data for the geometries, absorption spectra, and NTOs were generated using Chemcraft 1.8.

2.4. Solar cell preparation and characterization

All OSCs with a conventional device structure were fabricated onto detergent and solvent cleaned 150 nm thick ITO coated glass substrates (Colorado Concept). MoO₃ layers (5 nm) were deposited by thermal evaporation on the top of glass/ITO substrates in a vacuum chamber with base pressure of 10⁻⁸ Torr at a rate of 1 Å/s. The small molecule donor compounds **6a** and **6b** (21 mg of donor) were blended with either PC₆₀BM or PC₇₀BM (Nano-C) fullerene acceptor material (14 mg) in 1.5:1 weight ratio in 1 mL of a solvent mixture at 60 °C for 1 hour. The solvent mixture was prepared by mixing 19.92 mL of chlorobenzene with 80 µL of solvent additive 1,8-diiodooctane (Sigma-Aldrich). The 80 nm thick BHJ active layers were processed using spin coating (1400 rpm for 60 seconds). The thicknesses of the active layers were measured by a Dektak 6M profilometer. After the spin coating, the devices were subsequently annealed at 100 °C for 10 min to remove possible residual solvents. The device structures were completed with thermally evaporated 20 nm thick calcium (Ca) layers covered with 150 nm thick silver (Ag) layer acting as the cathode. The Ca and Ag layers were deposited via thermal evaporation in a vacuum chamber with base pressure of 10⁻⁸ Torr at rates of 3 Å/s and 6 Å/s, respectively. All processing steps were done inside the N₂ filled glove box. The active area of the OSC devices in all cases were 0.13 cm². Current density-voltage

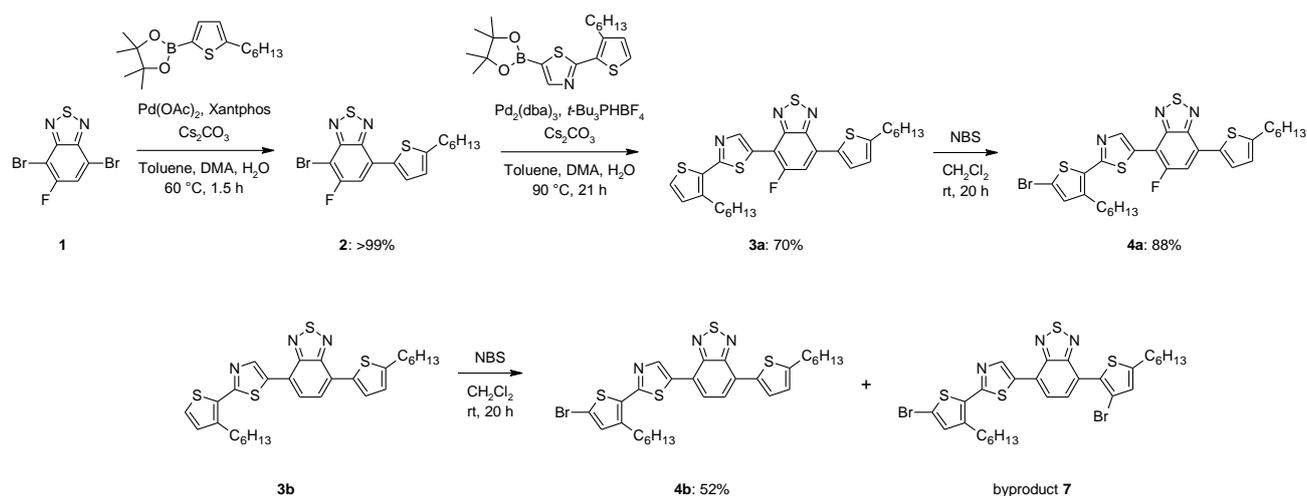
characteristics of the prepared OSCs were measured in dark and under simulated solar light, using a Keithley 2602A in combination with an Abet solar simulator, calibrated to produce 100 mW/cm² AM1.5G illumination. All device measurements were performed in a N₂ environment and at room temperature. For each sample, the current-voltage measurement data is presented as average of 12 cells with error bars (standard deviations) in Figures S4 and S5 in the Supplementary material. The results are reported as the values of the best cells in Figure 6.

3. Results and Discussion

3.1. Syntheses

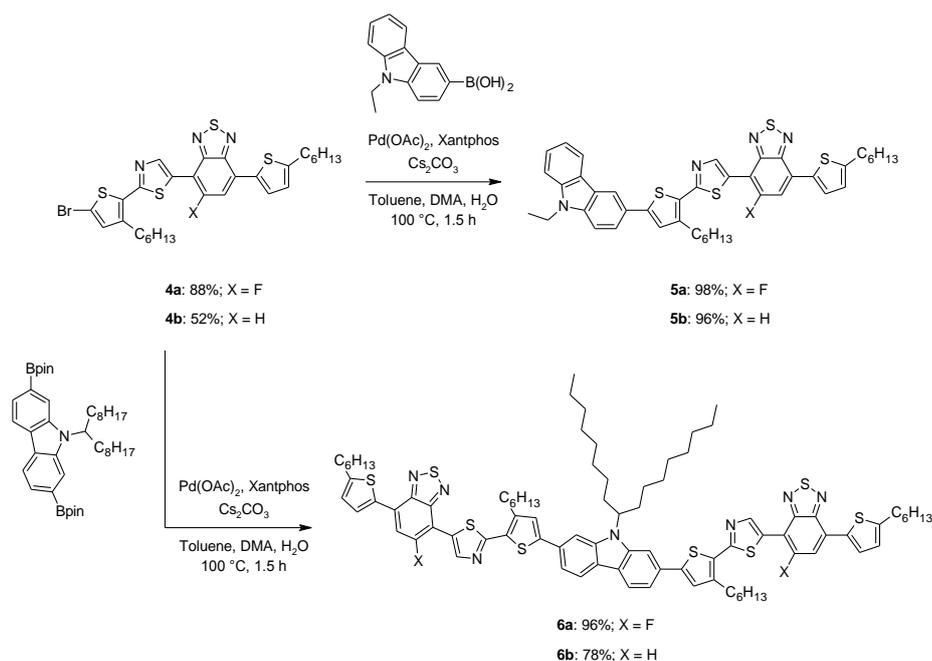
Compound **2** has previously been synthesized from compound **1** in 50% yield using Stille cross-coupling approach [37]. We found that a regioselective Suzuki-Miyaura cross-coupling between compound **1** and 5-hexyl-2-thiopheneboronic acid pinacol ester occurs in the presence of Pd(OAc)₂/Xantphos catalyst system. Moreover, the desired compound **2** was collected in quantitative yield (Scheme 1). The structure of compound **2** was confirmed by ¹H 1D and ¹H-¹H 2D NOE NMR experiments (see Supplementary material). At the next step, compound **2** was cross-coupled with 2-(3-hexylthiophen-2-yl)-5-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)thiazole [27] affording compound **3a** in low 35% yield in the presence of Pd(OAc)₂ as a palladium source. The yield was improved up to 70% by the replacement of Pd(OAc)₂ with Pd₂(dba)₃.

Compounds **3a** and **3b** [27] were brominated with *N*-bromosuccinimide (NBS) in CH₂Cl₂. Surprisingly, these two substrates showed quite different reactivity and selectivity in bromination. Compound **4a** was isolated in good 88% yield, whereas the reaction with compound **3b** gave a complicated mixture and the desired product **4b** could be isolated only in 52% yield. The bromination of compound **3b** also resulted in a dibrominated byproduct **7**.



Scheme 1. Syntheses of the small molecule donor material building blocks

At the final stage, compounds **4a** and **4b** were cross-coupled with 9-ethylcarbazole-3-boronic acid (Scheme 2). The resulting *D–A* type small molecule donor materials **5a** and **5b** were isolated in high 98 and 96% yields, respectively. The Suzuki-Miyaura cross-coupling reaction between 9-(9-heptadecanyl)-2,7-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)carbazole and compounds **4a** and **4b** gave the symmetrical *A–D–A* type small molecule donor materials **6a** and **6b** in 96 and 78% yields, respectively. The overall yields of fluoro substituted donor materials **5a** and **6a** were 34 and 33%, respectively, after seven reaction steps [38]. The total synthesis of compounds **5b** and **6b** were more difficult since, after six reaction steps, the overall yields were 21 and 17%, respectively [39].



Scheme 2. Syntheses of the *D-A* and *A-D-A* type small molecule donor materials

3.2. Spectroscopic and electrochemical properties

The experimental UV-vis absorption spectra of compounds **5a**, **5b**, **6a**, and **6b** in chloroform solution, and those of thin films, normalized at the S_0-S_1 transition absorption maximum, are shown in Figures 2 (A) and (B), respectively. The calculated time-dependent density functional theory (TDDFT) solution spectra are presented in Fig. S1. The main results retrieved from the spectra have been summarized in Table 1.

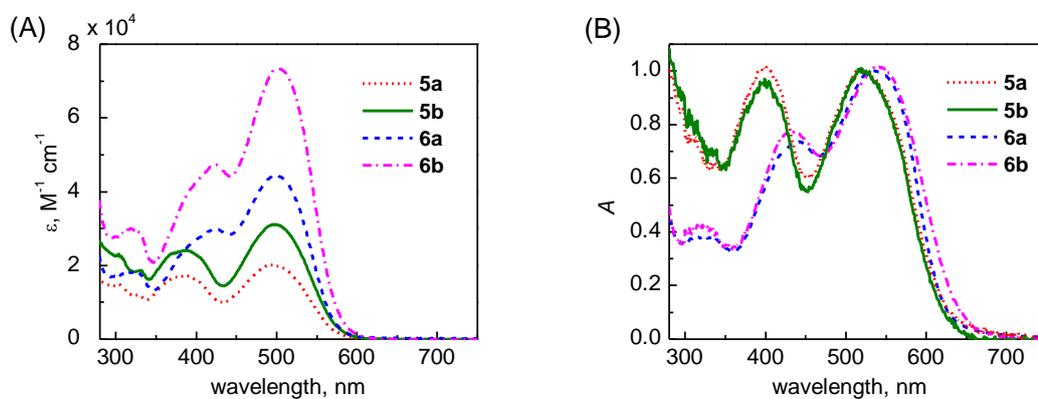


Fig. 2. (A): molar absorption coefficient (ϵ) of compounds **5a**, **5b**, **6a**, and **6b** in chloroform solution ($c = 7.5 \mu\text{M}$). (B): normalized absorption spectra of films of the target compounds.

Table 1Electrochemical and spectroscopic properties of compounds **3b**, **5a**, **5b**, **6a**, and **6b**

Compound	HOMO/LUMO (eV) ^a	E_{gap} (eV) ^a	HOMO/LUMO (eV) ^b	E_{gap} (eV) ^b	λ_{max} (nm) ^c	ε at λ_{max} (M ⁻¹ cm ⁻¹) ^c	λ_{max} (nm) ^d	$E_{\text{vert,max}}/\lambda^e$ (eV/nm)	f^f
3b ^g	-5.49/-3.13	2.36	-	-	475	35 300	486	-	-
5a	-5.29/-3.29	2.00	-5.00/-2.73 (-4.85/-2.63)	2.27 (2.22)	495	20 050	400; 520	2.61/475 (2.63/472)	1.40 (1.17)
5b	-5.25/-3.13	2.12	-5.00/-2.69 (-4.87/-2.56)	2.31 (2.31)	499	31 115	400; 520	2.58/481 (2.62/473)	1.41 (1.22)
6a	-5.20/-3.16	2.04	-5.01/-2.77 (-4.84/-2.69)	2.24 (2.15)	500	44 375	441; 534	2.59/479 (2.59/478)	2.82 (2.51)
6b	-5.28/-3.27	2.01	-5.01/-2.73 (-4.88/-2.63)	2.28 (2.25)	503	73 380	436; 541	2.56/485 (2.58/480)	2.79 (2.58)

^aDPV in dichloromethane. ^bCalculated at the DFT/B3LYP/6-31G** level of theory in chloroform and in vacuum (in parentheses). Wavelengths correspond to the absorption maxima in ^cchloroform solution (7.5x10⁻⁶ M) and in ^dfilm samples, spin-coated from chloroform solution. ^eIn chloroform and in vacuum (in parentheses); calculated at the TDDFT/ ω B97X-D/6-31G** level of theory using the B3LYP/6-31G** optimized geometries. The reported transition is for the one with the largest ^foscillator strength (f), i.e. absorption maximum, which corresponds to the S₀→S₁ transition for all compounds. ^fOscillator strengths calculated in chloroform and in vacuum (in parentheses). ^gFrom reference 27.

Both the experimental and theoretically calculated absorption spectra of the four molecules in chloroform solution have very similar features with two strong, distinct bands centred at around 400 nm (violet band) and 500 nm (cyan band). The former band corresponds to a π - π^* electronic transition of the chromophores. The latter band is attributed to intra-molecular charge transfer (ICT) from the *D* to the *A* moiety. This ICT is also observed theoretically in the presentation of the hole and electron natural transition orbitals (NTOs) of the main S₀→S₁ transition (with the highest oscillator strength) calculated with TDDFT, see Table 1 and the studies of the structural and charge transfer properties below. As can be seen from Fig. 2, for all the molecules the intensity of the cyan band is higher than that of the corresponding violet band. Furthermore, for compounds **5a**, **5b**, **6a**, and **6b**, the absorption maxima are practically overlapping. Clearly, the extension of molecular structure with another *A*- π unit as in compounds **6a** and **6b** has only a minor effect on the absorption range. The introduction of fluorine in the *A* units, with its well-known inductive effect, is responsible for the small blue-shifts of the cyan bands in **5a** and **6a** relative to the non-fluorinated compounds **5b** and **6b**. Interestingly,

the fluoro substituent decreased notably molar absorption coefficients (ϵ) of compounds **5a** and **6a** relative to the non-fluorinated counterparts **5b** and **6b**, respectively. This effect is opposite to the previous observation of fluoro substitution in *D-A-D* type small molecule electron donors [40]. Similarly, the TDDFT calculations predict small blue-shifts of ca. 6 nm for the fluorinated compounds relative to the non-fluorinated ones (Table 1, Fig. S1). Moreover, the calculated oscillator strength of the fluorinated **5a** in chloroform is slightly decreased compared to **5b**, whereas for the symmetric **6a** and **6b** the opposite is predicted. By comparing the absorption properties of compounds **3b** and **5b**, the effect of the Cz unit can be clearly seen: the cyan band of **5b** is red-shifted by 24 nm compared to the cyan band of **3b**. However, the molar absorption coefficients are nearly the same. This result is expected, since the biggest contribution to a high ϵ value comes from the effective conjugation between the fused BT unit and thiophene/thiazole units. The biggest difference in the absorption properties of unsymmetric compounds **5a** and **5b** compared to symmetric compounds **6a** and **6b** is that ϵ values of the symmetrical **6a** and **6b** are increased notably due to their extended molecular structures. The same trend is observed in the calculated TDDFT spectra (Fig. S1), i.e. the oscillator strengths of **6a** and **6b** are significantly larger than those of **5a** and **5b** (Table 1).

In solid state spectra, the cyan bands are significantly red-shifted with respect to the cyan bands of the solution spectra, due to chromophore-chromophore intermolecular interactions in solid state. This is most evident for compounds **6a** and **6b** due to their extended molecular structures. The same phenomenon can be observed by comparing the solid state absorptions of compounds **3b** and **5b**. The cyan band of **5b** is red-shifted (34 nm) compared to the cyan band of its building block **3b**. The Cz unit extends the molecular structure of **5b** with respect to **3b**, increasing the intermolecular interactions and, thus, causing even more pronounced red-shift of absorption in solid state than in solution. The fluoro substituent does not significantly affect the intensities and the wavelengths of cyan bands of compounds **5a** and **5b** are equal. The cyan band of **6a** is blue-shifted with respect to **6b** as in solution spectra by the fluoro substituent, while the violet band of fluorinated compound **6a**

is red-shifted relative to non-fluorinated compound **6b**. The TDDFT calculations in vacuum predict the same trends, namely blue-shifted spectra of the fluorinated **5a** and **6a** with respect to the non-fluorinated **5b** and **6b**. Additionally, the increased conjugation in the symmetric compounds **6a** and **6b** red-shifts their vacuum spectra with respect to the unsymmetric **5a** and **5b**. The fluoro substituent decreases somewhat more the calculated oscillator strengths in **5a** and **6a** compared to the solvent calculations.

The electrochemical properties, *i.e.* the energies of the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO), of **5a**, **5b**, **6a**, and **6b** were determined by differential pulse voltammetry (DPV) in dichloromethane solution. In fact, the voltammograms, *i.e.* changes in current-voltage curves, presented with respect to the ferrocene couple in Fig. 3, clearly show reversible oxidation and reduction processes [41]. They are presented in Table 1 together with the HOMO–LUMO gap energies (E_{gap}) and the calculated HOMO, LUMO, and gap energies in chloroform and vacuum. Electrochemical properties of the different compounds do not differ significantly. This is consistent with the observation that the absorption edges of all the compounds **5a**, **5b**, **6a**, and **6b** locate at the same wavelength region. The HOMO energies lie in the range of -5.20 to -5.29 eV and the LUMO energies lie between -3.13 and -3.29 eV. By contrast, when comparing the energy levels of **3b** and **5b**, the extension of molecular structure of **5b** by Cz unit raises the HOMO energy and simultaneously narrows the E_{gap} by 0.24 eV. The LUMO energy remains intact.

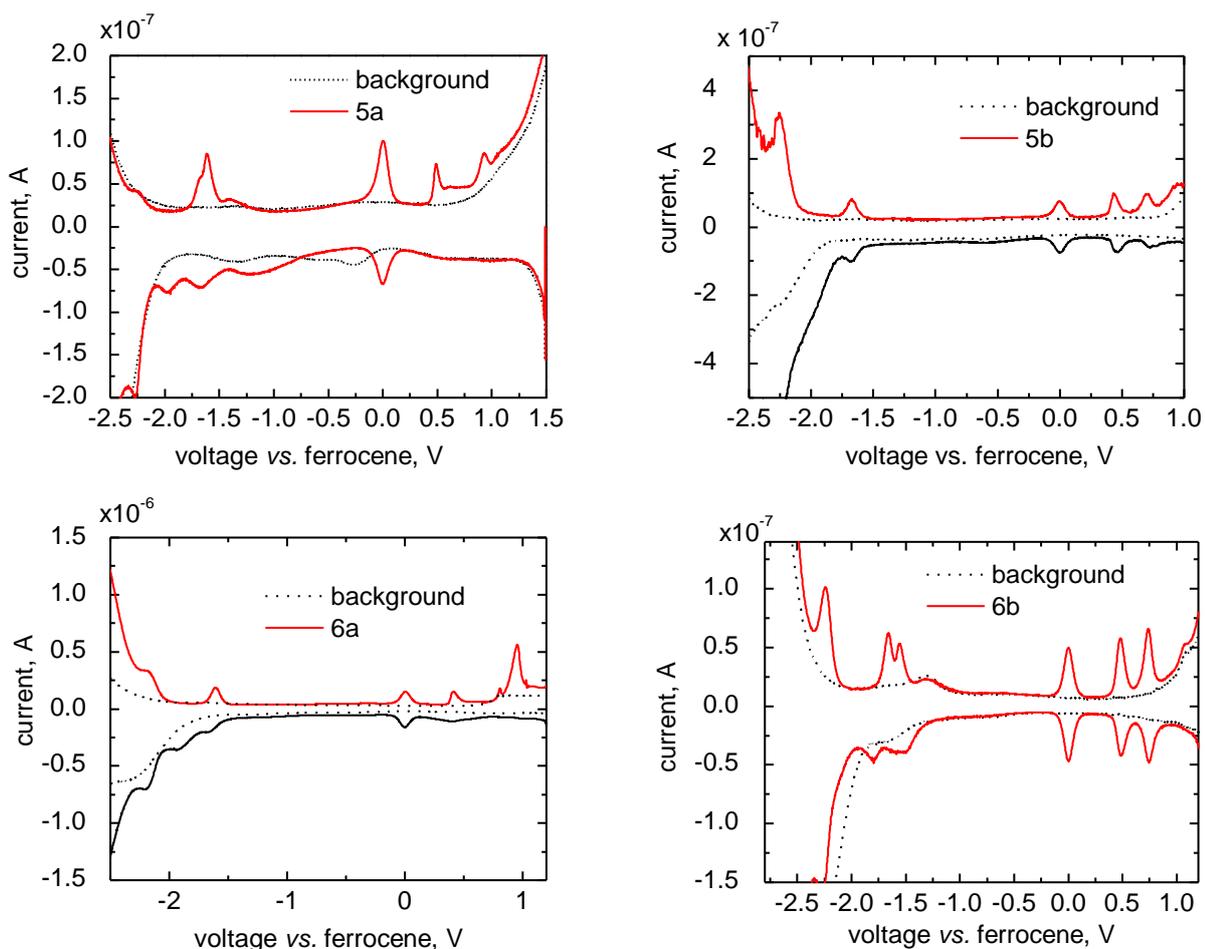


Fig. 3. DPV voltammograms of compounds **5a**, **5b**, **6a**, and **6b**.

3.3. Structural and charge transfer properties

The optimized ground-state geometries of the compounds are presented in Fig. 4. The adjoining thiophene, thiazole, and BT units are *anti* to each other, i.e. the sulfur atoms of the units are on the opposite sides of the molecules, in the energetically most preferable conformations of compounds **5a**, **5b**, **6a**, and **6b** (see Supplementary material, Fig. S2). In addition, thiophene and Cz are *syn* to each other in these structures, i.e. sulfur of thiophene and nitrogen of Cz are on the same side (α in Fig. 4). Moreover, conformational studies of **5a** suggest that the *anti*-orientation of thiophene with respect to Cz could also be possible, as the energy difference between these two conformations is only 0.17 kJ mol⁻¹. In the case of the symmetrical compound **6a**, the conformational

studies indicate that the *syn/syn*-orientation between thiophene and Cz units is more likely (by 1.25 kJ mol⁻¹) than the *syn/anti*-orientation, which may be due to the more favorable positioning of the side chains in the *syn/syn*-conformation. Both ¹H 1D and ¹H-¹H 2D NOE NMR data support (see Supplementary material) this prediction of the *syn/syn*-orientation at elevated temperatures (325–330 K) in chloroform. However, it is evident that also *syn/anti*-orientation between thiophene and Cz units exists for both compounds **6a** and **6b** at 297 K. On the grounds of the closest atom distances between the BT unit and the neighboring thiazole/thiophene π -bridge unit (the C–H \cdots N distance is ca. 2.3 Å and the C–H \cdots S distance is 2.7 Å), there is hydrogen bonding [42] between these adjoining units. This hydrogen bonding stabilizes the structure, and planarizes the molecular backbones of all compounds for the most part in both vacuum and chloroform (Fig. S2 and Table S1). However, the steric hindrance of hydrogen atoms causes torsional twists of ca. 24–27° between Cz and the neighboring thiophene π -bridge units (α in Fig. 4). Overall, the dihedrals are only slightly smaller when calculated in chloroform instead of vacuum. On the other hand, the size of the molecule does not influence the torsions of the molecular backbones, i.e. the dihedrals of the larger symmetrical compounds **6a** and **6b** are very close to those in the corresponding unsymmetrical compounds **5a** and **5b**, respectively. Moreover, the fluoro substituents of compounds **5a** and **6a** do not affect planarization.

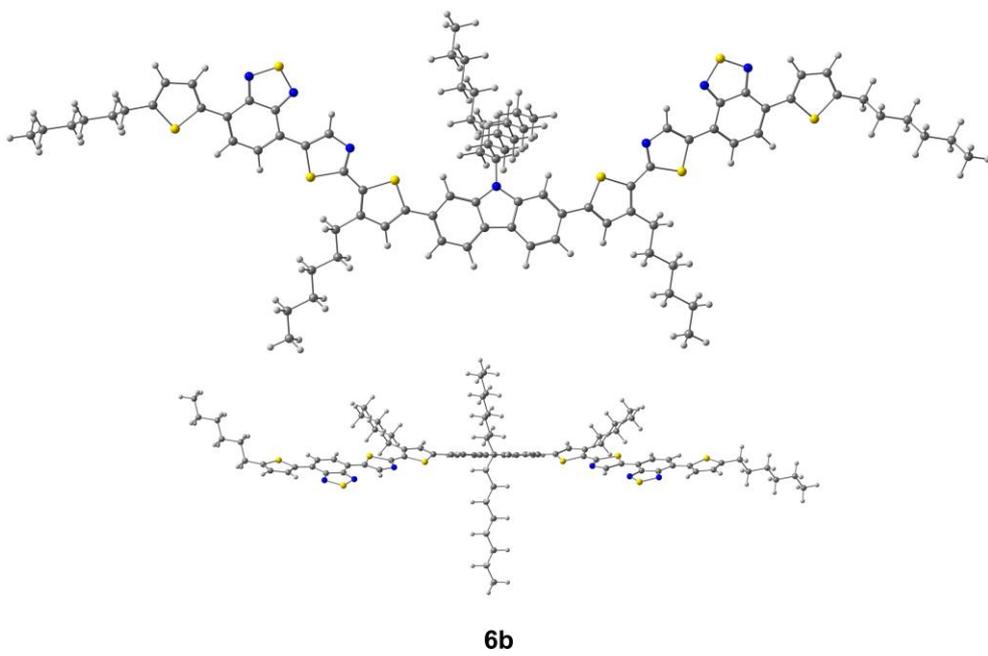
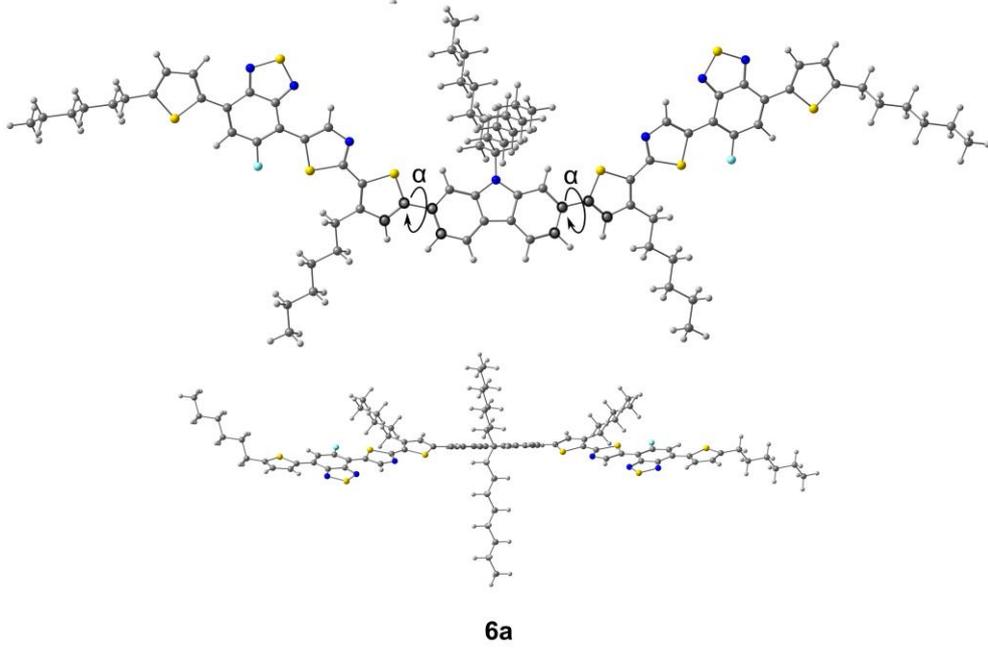
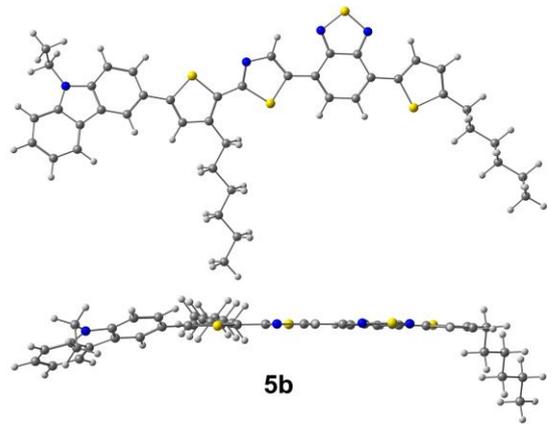
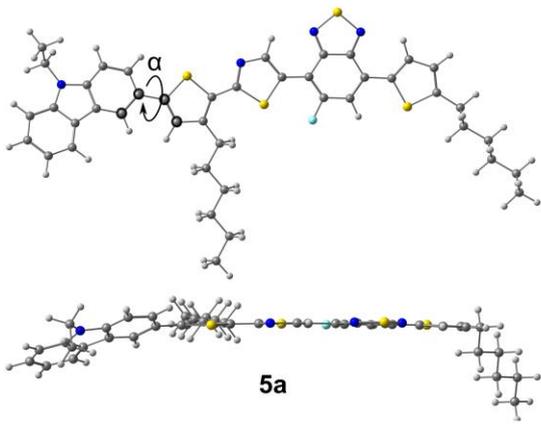


Fig. 4. Optimized ground-state geometries of compounds **5a**, **5b**, **6a**, and **6b** calculated in chloroform at the DFT/B3LYP/6-31G** level of theory. Dihedral angles, α , between Cz and thiophene in **5a** and **6a**, are defined the same way for compounds **5b** and **6b**.

The NTOs for the $S_0 \rightarrow S_1$ transitions, which correspond to the main absorption peak in the theoretical UV-Vis absorption spectra calculated with the ω B97X-D functional (see Supplementary material, Fig. S1), are represented in Fig. 5. The calculations predict one dominant NTO pair (λ_{NTO1} in Fig. 5) for the unsymmetrical *D-A* type compounds **5a** and **5b**, whereas two NTO pairs (λ_{NTO1} and λ_{NTO2} in the figure) with almost identical contributions (50% and 41%) are required to describe the $S_0 \rightarrow S_1$ transition of symmetrical *A-D-A* type compounds **6a** and **6b**. The corresponding NTOs of **5a** and **5b** are almost identical, the only difference being a small amount of charge density localized on fluoro substituent in **5a**. The same applies to **6a** and **6b**. Moreover, all the compounds have very similar ICT character: the holes delocalize more on thiophene/thiazole units and the six-membered ring of the BT units, while the electrons localize more on the thiazole and BT units. Strikingly, the Cz donor unit does not participate much to the charge distribution in all cases, which is most likely due to the larger torsional twists in the backbone around it. In the symmetrical compounds **6a** and **6b**, the charge density localizes more on the other end of the molecular backbone. The second NTO pairs of compounds **6a** and **6b** have the opposite charge density distributions compared to their first NTO pairs.

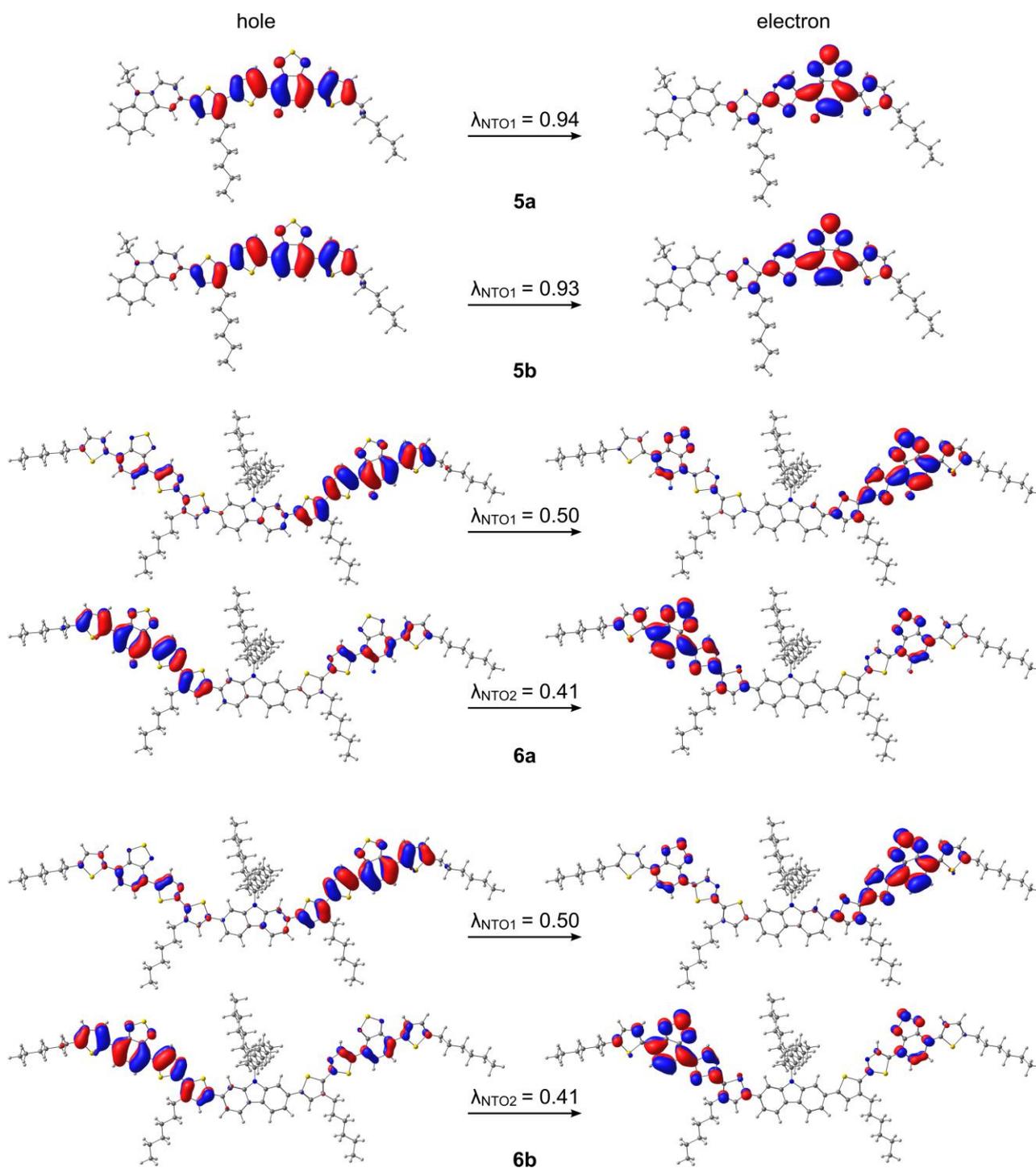


Fig. 5. NTOs for the $S_0 \rightarrow S_1$ transitions of compounds **5a**, **5b**, **6a**, and **6b** calculated at the TDDFT/ ω B97X-D/6-31G** level of theory in chloroform (isodensity contour = 0.025). The eigenvalue λ_{NTO} is the fraction of the hole–electron contribution to the given transition.

3.4. Photovoltaics

In order to test the potential of synthesized organic compounds as donor materials for optoelectronic applications, BHJ OSCs were prepared with conventional device structure: glass/ITO/ \sim 5 nm MoO₃/80 nm active layer/ \sim 20 nm Ca/ \sim 150 nm Ag. For the active layers, donor materials **6a** and **6b** were mixed with fullerene derivatives in 1.5:1 weight ratio, respectively. 80 nm thick films were produced in all cases using spin-coating. Compounds **5a** and **5b** were omitted due to limited film forming properties. Current-voltage measurements of the best single cells in Fig. 6 demonstrate the possibility to use compounds **6a** and **6b** as organic electron donor materials for both type of fullerene derivatives (PC₆₀BM and PC₇₀BM). The device performances in dark demonstrate good diode characteristic with low leakage current, as shown in Fig. 6(B) and (D). When devices were illuminated with 1 sun simulated light, very good V_{oc} (0.81–0.88 V) were obtained in all cases.

The best device performances were observed using PC₇₀BM as an electron acceptor as shown in Fig. 6(A) and (C). The device with compound **6a** as an electron donor gave the highest obtained PCE of 2.80% with V_{oc} of 0.88 V, short-circuit current (J_{sc}) of 5.80 mA cm⁻², and fill factor (FF) of 55%. Down the line, the fluoro substituted compound **6a** outperformed **6b** as a donor component, since the best observed PCE with compound **6b** as an electron donor component (of 80 nm thick active layer) was 2.30%, with V_{oc} of 0.86 V, J_{sc} of 5.54 mA cm⁻², and FF of 48%. Similar positive effect of the mono-fluoro substitution of BT unit has been previously demonstrated for $D-A-D$ [40] and more extended $D-A-D-A-D$ [43,44] type small molecule electron donors. One reason for the common positive contribution of the fluoro substituent can be the increased intermolecular interactions due to C–H \cdots F–C and S \cdots F–C interactions that can lead to improved hole mobilities [44].

The limited performances of the prepared solar cell devices are mainly due to relatively low FF and J_{sc} . The careful optimization of e.g. both solvent mixture and active layer composition would likely produce better morphologies leading to higher FF s. However, the detailed optimization of OSC

devices is not the scope of this paper. Moreover, the design of the *A–D–A* molecular structure needs further studies, e.g. the other *D* units should be tested to reduce the twist between the central *D* unit and π -bridge, and to find out how this affects the performance of the compound as donor component of BHJ layer. Based on the computational studies, it seems that the twist between the central *D* unit and π -bridge may prevent effective ICT process that can be the reason for the observed relatively low J_{sc} values.

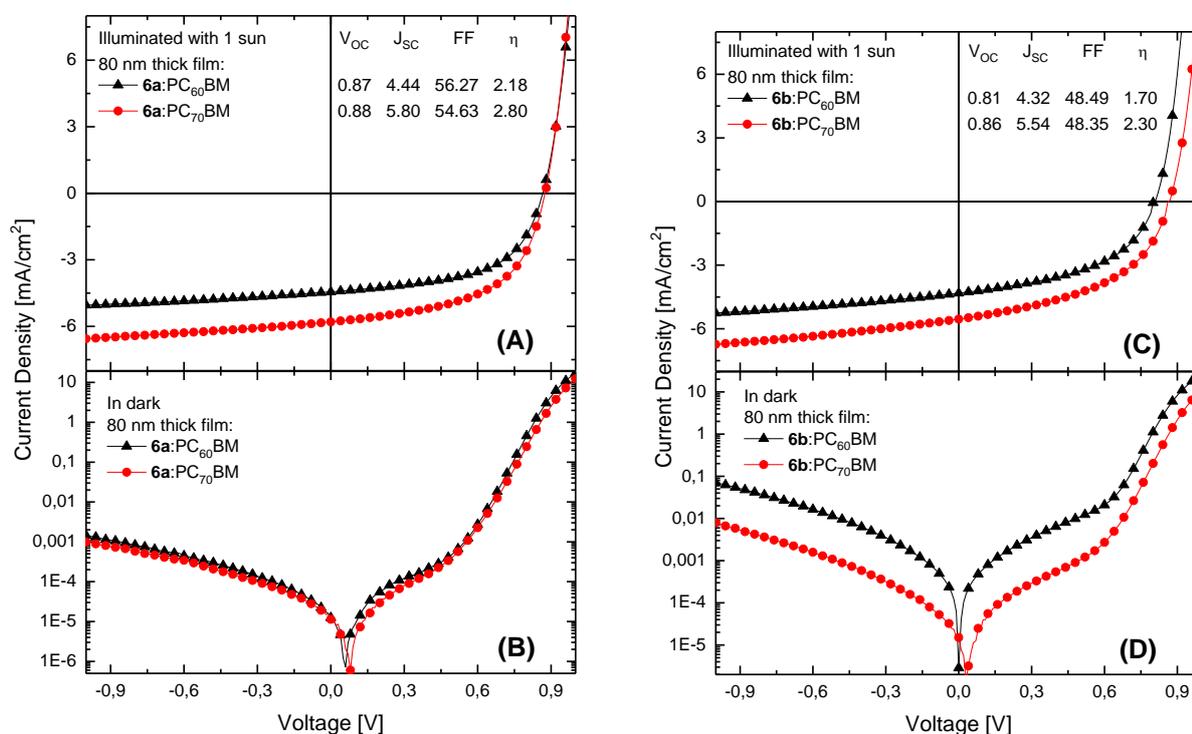


Fig. 6. Photovoltaic performance of the electron donor materials **6a** and **6b** mixed with different fullerene derivatives, (A) and (C) under illumination with 1 sun and (B) and (D) in dark.

4. Conclusions

Synthetic methods were developed to construct organic building blocks, which can be widely utilized in the syntheses of organic semiconducting materials. In this study, two *D–A* and two *A–D–*

A type small molecule materials were synthesized and characterized in detail. Finally, the A–D–A type materials were tested as donor components in BHJ-based OSC devices.

Results show clearly that both D–A and A–D–A type compounds have many similar material properties, e.g. the HOMO/LUMO energies of these two compound types differ only slightly. Also, the compounds have very similar nearly planar conformations. On the other hand, the symmetric A–D–A type materials **6a** and **6b** have several advantages over the unsymmetric D–A type materials **5a** and **5b**. For example, the extended molecular structure of the A–D–A type materials provides higher molar absorption coefficients and good film forming properties. These are crucial factors for efficient use of these materials in semiconducting applications. In solar cell tests, the fluoro substituent of compound **6a** is responsible for the enhanced device performance (especially due to the increased *FF*), when comparing compounds **6a** and **6b** as donors with both PC₆₀BM and PC₇₀BM acceptors. However, to fully optimize the structures of the A–D–A type materials, other central *D* units should be tested, to prevent the undesired twist of the molecular structures between the central *D* unit and the π -bridge. In this way, more effective conjugation and decreased E_{gap} values may be achieved. Small molecule donor materials with a narrower bandgap would cover a broader range of the solar spectrum and increase the total absorption leading to increased photo-current. All these factors are beneficial for the material performance as an active layer component in a BHJ-based OSC device.

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Supplementary Material

Supplementary data related to this article can be found at

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