Article

A-π-D-π-A-Based Small Molecules for OTFTs Containing Diketopyrrolopyrrole as Acceptor Units

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Abstract: A-π-D-π-A-based small molecules 6,6′−((thiophene-2,5-diylbis(ethyne-2,1-diyl))bis(thiophene-5,2-diyl))(2,5-bis(2-ethylhexyl)-3-(thiophene-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione) (TDPP-T) and 6,6′−(((2,3-dihydrothieno[3,4-b][1,4]dioxine)-5,7-diyl)bis(ethyne-2,1-diyl))bis(thiophene-5,2-diyl)(2,5-bis(2-ethylhexyl)-3-(thiophene-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione) (TDPP-EDOT) have been designed and synthesized. The diketopyrrolopyrrole acts as an electron acceptor, while the thiophene or 3,4-ethylenedioxythiophene acts as an electron donor. The donor–acceptor groups are connected by an ethynyl bridge to further enhance the conjugation. The optoelectronics, electrochemical, and thermal properties have been investigated. Organic thin film transistor (OTFT) devices prepared from TDPP-T and TDPP-EDOT have shown p-type mobility. In as cast films, TDPP-T and TDPP-EDOT have shown a hole mobility of 5.44 × 10⁻⁶ cm² V⁻¹ s⁻¹ and 4.13 × 10⁻⁶ cm² V⁻¹ s⁻¹, respectively. The increase in the mobility of TDPP-T and TDPP-EDOT OTFT devices was observed after annealing at 150 °C, after which the mobilities were 3.11 × 10⁻⁴ cm² V⁻¹ s⁻¹ and 2.63 × 10⁻⁴ cm² V⁻¹ s⁻¹, respectively.

Keywords: organic thin film transistors; hole mobility; small molecules; donor-acceptor; diketopyrrolopyrrole

1. Introduction

During the past decades, organic semiconductors have emerged as an alternative to inorganic semiconductors because of their superior advantages compared with their inorganic counterparts [1]. Among organic semiconductors, organic thin film transistors (OTFTs) are significant because of their many applications in fields like flexible displays [2], inexpensive radio frequency identification (RFID) tags [3], and sensors [4]. Exploiting the solution processable properties of OTFTs devices allows for them to be prepared easily at a low cost [5]. Tremendous work has been done by researchers on the synthesis of several small molecules and polymers constituting benzothiadiazole (BTD) [6–8], diketopyrrolopyrrole (DPP) [9–12], isoindigo (ii) [13–15], benzobisthiazole (BBT) [16], and arylene diimides [17–19], which have shown good mobility.

So far, OTFTs have been prepared from both polymers and oligomers. In general, polymers tend to achieve a high mobility for OTFTs. Nevertheless, the simple preparation and easy purification of small molecules makes them a better choice over polymers. So far, significant results have been reported on small molecule-based OTFTs, including
anthracene [20–22], naphthalene [23,24], thienothiophene [25,26], and fluorene [27], which have demonstrated good mobility.

OTFT performances can be improved by tuning the molecular orbitals of the organic molecules. The most useful and effective way of modifying the energy levels of molecular orbitals of organic molecules is achieved using a donor–acceptor (DA) strategy. The donor–acceptor molecules have a low energy gap between the HOMO and LUMO levels. The common electron acceptor molecules are benzothiadiazole [28], benzoselenadiazole [29], and diketopyrrolopyrrole [30], while the common electron donor molecules are thiophene [31], 3,4-ethylenedioxythiophene [32], and thienothiophene [33]. Although many acceptor molecules have been explored for OFETs, diketopyrrolopyrrole remains one of the most versatile and widely used molecules. Diketopyrrolopyrrole (DPP) has a planar conjugated bicyclic structure, which enables efficient intermolecular $\pi$-$\pi$ interactions, whereas the lactam part makes the DPP unit exhibit a strong electron-withdrawing effect. As for the donors in small molecule-based OTFTs, ethylene-based conjugated systems have gained decent attention from scientists and technologists [34]. The ethynylene bridge provides the diketopyrrolopyrroles coplanar to the central donor group, thus promoting extensive $\pi$-conjugation [35]. It also enhances the intramolecular charge transfer process [36]. In particular, for charge transfer purposes, the presence of unsaturated linkages (such as in ethylene and ethynylene) between the donor and acceptor units has been proven to be beneficial, as these linkages potentially reduce steric hindrance and enhance co-planarity along the molecular skeleton [37]. In this context, 3,4-ethylenedioxythiophene (EDOT) has a prominent position in the category of extended $\pi$-donors, because of its unique combination of strong electron donor properties and self-structuring effects related to the intramolecular non-covalent interactions between oxygen and sulfur [38]. These intramolecular interactions also exert a determining influence on the structure and electronic properties of donors incorporating EDOT units. However, so far, the polymeric counterpart of EDOT, i.e., poly (3, 4-ethylenedioxythiophene (PEDOT), has been extensively used for the purpose of solution processed organic electronics, while the application of EDOT is rare [39]. This motivated us to apply thiophene and 3,4-ethylenedioxythiophene as donors in small molecule-based OTFTs.

Herein, we report two novel A-$\pi$-D-$\pi$-A type conjugated small molecules, namely TDPP-T and TDPP-EDOT (Scheme 1). In the two compounds, diketopyrrolopyrrole acts as an electron acceptor, while thiophene and 3,4-ethylenedioxythiophene act as electron donors. Incorporation of the rigid ethynylene bridge can increase conjugation, thus the absorption bands can be broadened to a longer wavelength region, further narrowing the band gap. The synthesized compound was investigated for its optical, electrochemical, thermal, and OTFT properties. The TDPP-T and TDPP-EDOT were confirmed by NMR spectra. The Figures S1–S4 in the supporting information represent the NMR spectra of the TDPP-T and TDPP-EDOT.
Scheme 1. Schematic representation of A-π-D-π-A type conjugated small molecules, TDPP-T and TDPP-EDOT, for OTFTs.

2. Materials and Methods

2.1. Materials

Sodium, FeCl₃, 2-methyl-2-butanol, 2-ethyl hexyl bromide, Pd(PPh₃)₄, CuI, (i-Pr)₂NH, ethynyltrimethylsilane, N-bromosuccinimide, 3,4-ethylenedioxythiophene, and 2,5-dibromo-thiophene were purchased from Sigma-Aldrich. DMF and 2-carbonitrile thiophene were purchased from Alfa Aesar Co. Tetrahydrofuran, dichloromethane, toluene, methanol, ethyl acetate, hexane, CHCl₃, and potassium carbonate were purchased from Samchun Pure Chemicals. Tetrahydrofuran (THF), diethyl ether, toluene, and methylene chloride were used after distillation in the presence of sodium/benzophenone or calcium hydride under nitrogen gas.

2.2. Synthesis of Organic Compounds

2.2.1. Synthesis of 3,6-Di(thiophen-2-yl)pyrrolo[3,4-c]pyrrole-1,4(2H,5H)-dione (3)

A mixture of sodium (7.39 g, 321.6 mmol), FeCl₃ (0.31 g, 1.93 mmol), and dry 2-methyl-2-butanol (159 mL) was stirred at 120 °C for 3 h under an N₂ atmosphere. Then, 2-carbonitrile thiophene (2) (14.99 mL, 160.8 mmol) was added and stirred for 30 min. Then, diethyl succinate (1) (8.4 mL, 9.40 mmol) in dry 2-methyl-2-butanol (66 mL) was added. After stirring the reaction mixture at 120 °C for 24 h, the contents in the flask were cooled to 50 °C, then acetic acid (55 mL) was added. After that, the mixture was refluxed for 1 h, cooled back to room temperature, and poured into methanol (50 mL). The precipitate was filtered and washed with water and methanol to afford (3) a dark red solid (10.5 g, 54%), which was used without further purification.
2.2.2. Synthesis of 2,5-Bis(2-ethylhexyl)-3,6-di(thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione (4)

Compound 3 (21.75 g, 72.4 mmol) and potassium carbonate (40.01 g, 289.6 mmol) were refluxed in DMF (522 mL) at 120 °C for 1 h. Then, 2-ethyl hexyl bromide (48.936 g, 253.4 mmol) was added dropwise. The reaction mixture was stirred at 120 °C for 36 h and was then cooled to room temperature. CHCl₃ (100 mL) and water (100 mL) were added and the layers were separated using a separating funnel. The organic layer was washed with brine, and the solvent was evaporated using a rotatory evaporator. The resulting solid was subjected to column chromatography (silica, eluent hexane/CH₂Cl₂, v/v 1/1) to afford (4) a red solid (1.5 g, 16%).

\[ \text{H NMR (300 MHz, CDCl₃, ppm):} \delta 8.91 (dd, J = 3.9, 1.2 Hz, 2H), 7.64 (dd, J = 5.1, 1.2 Hz, 2H), 7.28 (t, 2H), 4.04 (dd, J = 7.8, 3.6 Hz, 4H), 1.90–1.84 (m, 2H), 1.40–1.24 (m, 28); \]

\[ \text{C NMR (75 MHz, CDCl₃, ppm):} \delta 161.8, 140.4, 130.5, 129.8, 128.4, 107.9, 45.9, 39.1, 30.2, 28.4, 23.5, 23.1, 14.0, 10.5. \]

2.2.3. Synthesis of 3-(5-Bromothiophen-2-yl)-2,5-bis(2-ethylhexyl)-6-(thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione (5)

To a compound 4 (2.5 g, 4.76 mmol) in chloroform (60 mL), NBS (0.68 g, 3.81 mmol) was added in small portions at room temperature and stirred the reaction mixture for overnight. Then water was added, and the reaction mixture was extracted three times with methylene chloride (50 mL). The collected organic layers were dried over MgSO₄, filtered, and the solvent was evaporated under reduced pressure. The crude residue was purified by silica gel chromatography with methylene chloride/n-hexane, (1:1, v/v) as eluent to give 5 as a red solid. (0.8 g, 27%).

\[ \text{H NMR (300 MHz, CDCl₃, ppm):} \delta 8.92 (dd, J = 3.9, 1.2 Hz, 2H), 7.64 (dd, J = 4.8, 1.2 Hz, 2H), 7.29 (dd, J = 5.1, 3.9 Hz, 2H), 4.06–4.02 (m, 4H), 1.89–1.86 (m, 2H), 1.42–1.19 (m, 20H), 0.91–0.84 (m, 14H); \]

\[ \text{C NMR (75 MHz, CDCl₃, ppm):} \delta 161.6, 161.4, 140.9, 138.9, 135.6, 135.1, 131.4, 131.3, 130.9, 129.8, 128.5, 108.1, 107.8, 39.1, 39.1, 30.2, 28.3, 23.6, 23.5, 23.1, 14.0, 10.5. \]

2.2.4. Synthesis of 2,5-Bis(2-ethylhexyl)-3-(thiophen-2-yl)-6-(5-((trimethylsilyl)ethynyl)thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione (6)

Compound 5 (0.5 g, 0.83 mmol), Pd(PPh₃)₄ (19.2 mg, 0.0166 mmol), and CuI (0.9 mg, 0.005 mmol) were added to a dried two-neck flask. Degassed anhydrous toluene (20 mL) and (i-Pr)₂NH (4.16 mL) were added, followed by the addition of ethynyltrimethylsilane (163 mg, 1.66 mmol). Then, the reaction mixture was stirred under a nitrogen atmosphere at 90 °C overnight. After cooling to room temperature, water was added, and the mixture was extracted three times with ethyl acetate. The organic layer was dried over MgSO₄ and was filtered. The solvent was evaporated under reduced pressure to afford compound 6 (557 mg) in a 90% yield. The crude product was used for the next step without further purification.

\[ \text{H NMR (300 MHz, CDCl₃, ppm):} \delta 8.84 (dd, J = 3.9, 0.9 Hz, 1H), 8.71 (d, J = 4.2 Hz, 1H), 7.55 (dd, J = 5.1, 0.9 Hz, 1H), 7.23 (d, J = 4.2, 1H), 7.18 (t, J = 4.2 Hz, 1H), 3.95–3.88 (m, 4H), 1.78–1.76 (m, 2H), 1.30–1.14 (m, 18H), 0.91–0.84 (m, 14H); \]

\[ \text{C NMR (75 MHz, CDCl₃, ppm):} \delta 161.9, 161.8, 141.0, 139.5, 135.9, 135.2, 133.8, 131.1, 130.8, 130.0, 128.9, 128.8, 128.3, 109.1, 108.3, 104.2, 97.0, 46.3, 46.2, 39.3, 30.5, 30.4, 28.6, 23.8, 23.3, 14.3, 10.7. \]

2.2.5. Synthesis of 2,5-Bis(2-ethylhexyl)-3-(5-ethynylthiophen-2-yl)-6-(thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-dione (7)

K₂CO₃ (173 mg, 1.25 mmol) was added to a stirred solution of compound 6 (550 mg, 1 mmol) in CH₂Cl₂ (44 mL) at room temperature, and the reaction mixture was further stirred for 30 min. Then, 10 mL of water was added, and the mixture was extracted three times with ethyl acetate. The organic layer was dried over MgSO₄ and was filtered. The solvent was evaporated under reduced pressure. The crude product was purified by column chromatography with CH₂Cl₂/n-hexane (v/v, 1:3) as an eluent to afford compound 7 (417 mg) in a 95% yield.

\[ \text{H NMR (400 MHz, CDCl₃, ppm):} \delta 8.93 (d, J = 4 Hz, 1H), \]
8.80 (d, J = 4 Hz, 1H), 7.65 (d, J = 4 Hz, 1H), 7.38 (d, J = 4 Hz, 1H), 7.29 (d, J = 4 Hz, 1H),
4.04–3.97 (m, 4H), 3.58 (s, 1H), 1.86 (br, 2H), 1.38–1.32 (m, 16H), 0.90–0.86 (m, 12H); 13C
NMR (75 MHz, CDCl3, ppm): δ 161.7, 161.5, 141.1, 139.0, 135.8, 134.7, 134.0, 131.0, 130.9,
129.9, 129.7, 128.5, 126.6, 109.0, 108.0, 85.2, 46.0, 45.9, 39.2, 39.1, 31.9, 30.2, 29.7, 29.3,
28.3, 27.2, 25.3, 23.1, 22.7, 14.1, 14.0, 10.5.

2.2.6. Synthesis of 5,7-Dibromo-2,3-dihydrothieno[3,4-b][1,4]dioxine (9)

3,4-Ethynedioxythiophene (8) (1.5 g, 10.55 mmol) was dissolved in 32 mL of DMF.
N-Bromosuccinimide (3.94 g, 22.155 mmol) was dissolved in 32 mL DMF and was added
to the reaction mixture dropwise at 0 °C. The solution was stirred at room temperature for
4 h under the exclusion of light. After the completion of the reaction, water was added and
filtered. The precipitate was washed with water and the resulting precipitate was dried
under xzvacuum to afford compound 9 (2.1 g, 66%) (Scheme 2). 1H NMR (300 MHz,
CDCl3, ppm): δ 4.2 (s, 2H); 13C NMR (75 MHz, CDCl3, ppm): δ 139.7, 85.6, 65.0.

2.2.7. Synthesis of 6,6'-((Thiophene-2,5-diylbis(ethyne-2,1-diyl))bis(thiophene-5,2-
diyl))bis(2,5-bis(2-ethylhexyl)-3-(thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-
dione) (TDPP-T)

Compound 7 (0.8 g, 1.45 mmol), Pd(PPh3)4 (34 mg, 0.029 mmol), and CuI (1.7 mg,
0.0087 mmol) were loaded into a dried two-neck flask. Degassed anhydrous toluene (29 mL)
and (i-Pr)2NH (7 mL) were added, followed by the addition of 2,5-dibromothiophene (10)
(140 mg, 0.58 mmol). Then, the reaction mixture was heated to 90 °C and stirred overnight.
After cooling to room temperature, water was added and the mixture was extracted three
times with ethyl acetate. The organic phase was dried over MgSO4 and filtered. Then,
the filtrate was concentrated under reduced pressure. The crude product was purified via
column chromatography with CH2Cl2/n-hexane (v/v, 2:1) as an eluent to afford
TDPP-T (397 mg) in a 58% yield (Scheme 2). 1H NMR (300 MHz, CDCl3, ppm): δ 8.96 (dd, J = 3.9,
1.2 Hz, 2H), 8.88 (d, J = 4.2 Hz, 2H), 7.67 (dd, J = 5.1, 1.2 Hz, 2H), 7.42 (d, J = 4.2 Hz, 2H),
7.30 (dd, J = 5.1, 3.9 Hz, 2H), 7.26 (s, 2H), 4.07–4.02 (m, 8H), 1.91–1.87 (m, 4H), 1.43–1.26
(m, 32H), 0.95–0.88 (m, 24H); 13C NMR (75 MHz, CDCl3, ppm): δ 161.6, 161.5, 143.9, 140.8, 139.1,
135.7, 135.2, 133.4, 132.8, 131.4, 131.0, 129.8, 128.5, 127.1, 124.7, 109.0, 108.0, 90.1, 87.9, 46.1,
46.0, 39.2, 39.1, 30.2, 30.1, 28.4, 23.5, 23.1, 14.1, 14.0, 10.5.

2.2.8. Synthesis of 6,6'-'(((2,3-Dihydrothieno[3,4-b][1,4]dioxine-5,7-diyl))bis(ethyne-2,1-diyli)
bis(thiophene-5,2-diyl))bis(2,5-bis(2-ethylhexyl)-3-(thiophen-2-yl)-2,5-dihydropyrrolo[3,4-c]pyrrole-1,4-
dione) (TDPP-EDOT)

Compound 7 (0.65 g, 1.175 mmol), Pd(PPh3)4 (27 mg, 0.029 mmol), and CuI (1.3 mg,
0.007 mmol) were loaded into a dried two-neck flask. Degassed anhydrous toluene (23 mL)
and (i-Pr)2NH (6 mL) were added, followed by the addition of compound 9 (0.14 g, 0.47 mmol).
Then, the reaction mixture was heated to 90 °C and stirred overnight. After cooling to room
temperature, water was added and the mixture was extracted three times with ethyl acetate.
The organic phase was dried over MgSO4 and was filtered. Then, the filtrate was concentrated
under reduced pressure. The crude product was purified via column chromatography with
CH2Cl2/n-hexane (v/v, 2:1) as an eluent to afford TDPP-EDOT (0.35 g) in a 60% yield. 1H
NMR (300 MHz, CDCl3, ppm): δ 8.95 (dd, J = 3.9, 1.2 Hz, 2H), 8.89 (d, J = 4.2 Hz, 2H), 7.66
(dd, J = 5.1, 1.2 Hz, 2H), 7.41 (d, 4.2Hz, 2H), 7.23 (dd, J = 5.1, 3.9 Hz, 2H), 4.38 (s, 4H), 4.07–4.01
(m, 8H), 1.91–1.87 (m, 4H), 1.42–1.23 (m, 32H), 0.94–0.85 (m, 24H); 13C NMR (75 MHz, CDCl3,
ppm): δ 161.7, 161.6, 143.9, 140.8, 139.1, 135.7, 135.3, 133.3, 131.2, 130.9, 129.8, 128.5, 127.4, 108.9,
108.1, 100.3, 90.4, 87.8, 64.9, 46.1, 46.0, 39.1, 39.1, 30.2, 30.1, 28.4, 23.5, 23.1, 14.0, 10.5.
2.3. Measurements

The $^1$H and $^{13}$C NMR spectra were measured using an NMR spectrometer (Bruker DRX-300, Berlin, Germany). The values of the chemical shift were stated in $\delta$ units (ppm). Mass spectral investigation was conducted on a JMS-700, JEOL. The absorption studies were attained by using a UV–VIS spectrophotometer (Shimadzu, UV-1065PC, Tokyo, Japan). Cyclic voltammogram examinations were conducted using an epsilon $E_3$ at room tempera-

Scheme 2. Synthesis of TDPP-T and TDPP-EDOT.
ture (RT) in a 0.1M tetrabutylammonium perchlorite solution in chloroform under an N₂ atmosphere at a 50 mV/s scan rate. The Ag/AgCl and Pt wire served as the reference and counter electrodes, respectively. The thermal properties were examined using a thermogravimetric analysis (TGA) and differential scanning calorimeter (DSC) (Mettler Toledo AG, Analytical, Schwerzenbach, Switzerland). A DSC analysis was conducted under an N₂ atmosphere, at temperature ranges from 0–400 °C at a heating rate of 10 °C/min. The thermogravimetric analysis (TGA) was measured under an N₂ atmosphere, at a heating rate from room temperature to 800 °C, at a of 10 °C/min using a thermogravimetric analyzer. The OFET top-contact and bottom-gate were made-up on a common gate on highly n-doped silicon with a thick, thermally grown 300 nm SiO₂ dielectric layer. Further modification of the dielectric SiO₂ layer was carried out using a self-assembled octyltrichlorosilane (OTS) monolayer. The solution comprising the semiconductor was spin-coated at 2000 rpm from 0.5–1 wt.% in chloroform to attain the thin film. Gold source and drain electrodes were evaporated on top of the semiconductor layer. Entire the investigation procedure was carried out by a home-built lab-view program combined with Keithley electronic units in a nitrogen atmosphere.

3. Results and Discussions
3.1. Optical and Electrochemical Properties of the TDPP-T and TDPP-EDOT

The optical properties of TDPP-T and TDPP-EDOT were investigated using UV–VIS absorption spectroscopy. The UV–VIS absorption spectra of TDPP-T and TDPP-EDOT were investigated using UV–VIS absorption spectroscopy. The UV–VIS absorption spectra of TDPP-T and TDPP-EDOT in a chloroform solution and in thin films are shown in Figure 1. The resulting data are summarized in Table 1. The UV–VIS absorption spectra of the two compounds exhibited two absorption bands in both the solution and film states. The higher wavelength band corresponds to intramolecular charge transfer (ICT) between donor and acceptor groups, which is typical in donor–acceptor-based compounds. In solution, TDPP-T showed absorption peaks at 391 and 591 nm, whereas TDPP-EDOT showed at 410 and 594 nm. In the film state, the absorption peaks for TDPP-T and TDPP-EDOT were observed at 376, 607, and 630 nm, and 370, 415, and 663 nm, respectively. In both compounds the ICT band in the film state were broad and red-shifted when compared with the solution state because of the intermolecular interactions in the film state. The optical band gap energy calculated for TDPP-T and TDPP-EDOT from the thin film state was 1.73 eV and 1.70 eV, respectively. In TDPP-EDOT, 3,4-ethylenedioxythiophene had a strong electron-releasing group compared with the thiophene in TDPP-T [40]. Because of the enhanced electron-releasing effect, it increased the HOMO level and resulted in a reduction of the band gap. The donor–acceptor character contributed to a narrowing of the band gap energy between the HOMO and LOMO levels.

![Figure 1](image-url)

Figure 1. UV–VIS absorption spectra of (a) TDPP-T and (b) TDPP-EDOT in chloroform solution and thin films.
To investigate the electrochemical properties of TDPP-T and TDPP-EDOT, cyclic voltammetry (CV) measurements were employed, and the results are summarized in Table 1. The cyclic voltammetry (CV) of TDPP-T and TDPP-EDOT are shown in Figure 2. Both TDPP-T and TDPP-EDOT showed oxidation and reduction waves. The HOMO energy levels of TDPP-T and TDPP-EDOT were calculated from the onset oxidation potentials to be $-5.44 \text{ eV}$ and $-5.31 \text{ eV}$, respectively. In TDPP-T, the donor thiophene group was less electron releasing compared with 3,4-ethylenedioxythiophene in TDPP-EDOT, and, as a result, TDPP-T had a deeper HOMO level. The LUMO energy levels were evaluated to be $-3.71 \text{ eV}$ and $-3.61 \text{ eV}$, respectively, by using the HOMO values and optical band gaps (LUMO = $E_{\text{opt}} - E_{\text{HOMO}}$).

Figure 2. Cyclic voltammogram curves of (a) TDPP-T and (b) TDPP-EDOT.

3.2. Thermal Properties of the TDPP-T and TDPP-EDOT

The thermal properties of TDPP-T and TDPP-EDOT were investigated using thermo gravimetric analysis (TGA) and differential scanning calorimetry (DSC). The TGA and DSC curves are shown in Figure 3. The thermal properties are summarized in Table 1. The thermal properties of the materials used as the organic thin film transistor have a major influence on the stability and lifetime of the resulting device. The decomposition temperatures (5% weight loss, $T_d$) for compounds TDPP-T and TDPP-EDOT were observed at 395 and 383 °C, respectively, indicating that the compounds were thermally stable and could be used for OTFTs. The thermal decomposition temperature at 5% wt. loss for TDPP-T was further compared with TDPP-EDOT. Hence, TDPP-T was more thermally stable than the TDPP-EDOT. In the DSC analysis, the melting temperatures ($T_m$) of TDPP-T and TDPP-EDOT were observed at 201 and 211 °C, respectively.

### Table 1. Optical, electrochemical, and thermal properties of TDPP-T and TDPP-EDOT.

| Compound   | $\lambda_{\text{abs}/\text{nm}}$ | $E_g$ a (eV) | HOMO (eV) b | LUMO (eV) c | $T_d$ (°C) d | $T_m$ (°C) e |
|------------|-------------------------------|-------------|-------------|-------------|-------------|-------------|
| TDPP-T     | 391, 591                      | 376, 607,630| 1.73        | $-5.44$     | 395         | 201         |
| TDPP-EDOT  | 410, 594                      | 370, 415,663| 1.70        | $-5.31$     | 383         | 211         |

a Calculated from the absorption edge wavelength. b Calculated from the onset of the oxidation potential. c Calculated from the HOMO level-band gap. d Decomposition temperature at 5% weight loss. e $T_m$, endothermic melting point.
Figure 3. TGA curves of (a) TDPP-T and (b) TDPP-EDOT, DSC curves of (c) TDPP-T and (d) TDPP-EDOT.

3.3. OTFT Characterization of the TDPP-T and TDPP-EDOT

OTFT devices were fabricated using bottom gate and top contact (BGTC) configuration. The OTFT films were prepared using the SiO$_2$ insulator while octyltrichlorosilane (OTS) was used for the surface treatment. Active layers were deposited by spin-coating a solution of semiconductors (5 mg mL$^{-1}$ in CHCl$_3$) at 1500 rpm. The thickness of the thin film was 30 nm. The source and drain electrodes were deposited by thermal evaporation of Au through a shadow mask. The Au thickness was 100 nm. The hole mobility ($\mu$) was determined using the following equation in the saturation regime: $I_{ds} = (W/C_i) \times \mu \times (V_{gs} - V_T)^2$, where $C_i$ is the capacitance per unit area of the SiO$_2$ dielectric ($C_i = 15$ nF cm$^{-2}$), $I_{ds}$ is the drain-source current, $V_{gs}$ is the gate voltage, and $V_T$ is the threshold voltage. The OTFT devices were tested using a Keithley 4200 semiconductor characterization system under a nitrogen atmosphere. The voltage–current graphs of the OTFT devices are shown in Figure 4. Both TDPP-T and TDPP-EDOT showed a p-type mobility. In the as-cast film, TDPP-T and TDPP-EDOT showed a hole mobility of $5.44 \times 10^{-6}$ cm$^2$ V$^{-1}$ s$^{-1}$ and $4.13 \times 10^{-6}$ cm$^2$ V$^{-1}$ s$^{-1}$, respectively. After thermal annealing, the mobilities increased because the crystallinity was enhanced. The optimal annealing temperatures for TDPP-T and TDPP-EDOT were 150 °C. At this temperature, the devices showed a hole mobility of $3.11 \times 10^{-4}$ cm$^2$ V$^{-1}$ s$^{-1}$ and $2.63 \times 10^{-4}$ cm$^2$ V$^{-1}$ s$^{-1}$, respectively. The large threshold voltage of 42.6 V for the as-cast TDPP-EDOT devices is due to the poor interface conditions. Thermal annealing causes the modification the Au/SiO$_2$ interface, which further leads to a reduction of the threshold voltage [41].
Table 2. OTFT device performance of TDPP-T and TDPP-EDOT.

| µ_AVG (cm²/Vs) | V_TH (V) | On/Off Ratio |
|----------------|----------|--------------|
| TDPP-T         | TDPP-EDOT| TDPP-T       | TDPP-EDOT |
| As cast        | (5.44 ± 0.11) × 10⁻⁶ | (4.13 ± 0.16) × 10⁻⁶ | -5.0    | 42.6    | 13.2    | 3.5     |
| Ann 100 °C     | (3.61 ± 0.08) × 10⁻⁵ | (1.26 ± 0.06) × 10⁻⁵ | -1.5    | -2.5    | 23.1    | 73.4    |
| Ann 150 °C     | (3.11 ± 0.08) × 10⁻⁴ | (2.63 ± 0.12) × 10⁻⁴ | -11.5   | 1.4     | 326     | 1060    |

Figure 4. OTFT field-effect transistor characteristics. Output curves of (a) TDPP-T and (c) TDPP-EDOT. Transfer curves of (b) TDPP-T and (d) TDPP-EDOT.

The crystallinity and molecular ordering of thin films were investigated using X-ray diffraction (2D-GIXRD) patterns. The 2D-GIXRD patterns of TDPP-T and TDPP-EDOT are shown in Figures 5 and 6. The 2D-GIXRD patterns of TDPP-T and TDPP-EDOT were analyzed for the films as cast at 150 °C temperatures. In both compounds, TDPP-T and TDPP-EDOT, compared with the as cast film, thermal annealing at 150 °C increased the crystallinity, thus leading to an increase in mobility. The low mobilities were due to the poor crystallinity or low morphological orientation of the organic molecules. The mobility can be improved by using additives [42–45]. The organic molecules blended with additives cause phase segregation and increased crystallinity. Thus, the blended films will give allowance for improved OTFT performances.
Figure 5. Two-dimensional grazing incidence X-ray diffraction images of TDPP-T (a) without annealing and (b) with being annealed at 150 °C, corresponding line-cuts along the (c) q_{xy} and (d) q_{z} plane.

Figure 6. Two-dimensional grazing incidence X-ray diffraction images of TDPP-EDOT (a) without annealing and (b) with being annealed at 150 °C, corresponding line-cuts along the (c) q_{xy} and (d) q_{z} plane.
4. Conclusions

The donor–acceptor-based small molecule, TDPP-T and TDPP-EDOT were designed and synthesized. The band gap energy between the HOMO and LUMO was decreased due to the donor–acceptor interactions in the molecule. The alkyl chains promoted the good solubility of the compound in common organic solvents. TDPP-T- and TDPP-EDOT-based OTFT devices after thermal annealing at 150 °C have shown hole mobility of 3.11 x 10^{-4} cm^2 V^{-1} s^{-1} and 2.63 x 10^{-4} cm^2 V^{-1} s^{-1}, respectively.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/10.3390/mi12070817/s1. Figure S1. 1H NMR (300 MHz, CDCl3) spectrum of compound TDPP-T. Figure S2. 13C NMR (75 MHz, CDCl3) spectrum of compound TDPP-T. Figure S3. 1H NMR (300 MHz, CDCl3) spectrum of compound TDPP-EDOT. Figure S4. 13C NMR (75 MHz, CDCl3) spectrum of compound TDPP-EDOT.

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