Diacenopentalene dicarboximides as new n-type organic semiconductors for field-effect transistors†

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N,N'-Dihexyl-dibenzo[ghi]pentalene dicarboximide (DBPDI) and N,N'-dioctyl-dinaphthopentalene dicarboximide (DNPD) were successfully synthesized as new n-type semiconducting materials. They have good solubility in common organic solvents. DBPDI and DNPD are thermally stable, with decomposition temperatures at 419 and 460 °C, respectively, based on thermogravimetric analysis, and melt at 354 and 387 °C as measured by differential scanning calorimetry. Their optical and electrochemical properties were studied by UV-vis absorption and cyclic voltammetry measurements. Time-dependent density functional theory calculations (TD-DFT) were used to explain their unique electronic absorption spectra. DBPDI and DNPD have relatively low-lying LUMO energy levels at −3.76 eV and −3.45 eV and HOMO energy levels at −5.84 eV and −5.72 eV, respectively. Their application in organic field-effect transistors (FETs) was investigated. Both DBPDI and DNPD showed n-type field-effect transistor behavior. The DBPDI device obtained by solution-processing technique displayed an average electron mobility of up to 0.06 cm² V⁻¹ s⁻¹ with an I_on/I_off ratio of 2.9 × 10⁶.

Introduction

In the last decade, interest in the synthesis of cyclic π-conjugated systems incorporated with an antiaromatic unit has increased. Pentalene, as one of the important antiaromatic systems, has attracted much attention.¹ Parent pentalene, with its 8π electron framework, is unstable and easily dimerizes above −196 °C. Dibenzopentalene (DBP) has been synthesized by different methods, and it is a fairly stable compound with a planar structure and 4ππ electron periphery.² Incorporation of the antiaromatic pentalene unit into the acene framework is expected to be an efficient approach to stabilizing the acene moieties due to the intramolecular donor–acceptor interactions. As expected, the extended dinaphthopentalene (DNP)²,³ and dianthracenopentalene (DAP)⁴ derivatives were reported, and they can be used as p-type organic semiconductors with good stability. Other pentalene-containing molecules have also been studied for their applications in organic electronics.⁵ The pentalene unit can be regarded as a weak electron-withdrawing moiety that can stabilize the electron-rich moieties such as acenes but cannot convert them into n-type materials such as DNP,⁶ DAP,⁷ DBP-based polymers,⁸ pyrrole-fused DNP,⁹ and thienoacenefused pentalenes,¹⁰ which all show hole transport properties. To prepare n-type pentalene derivatives, other electron-deficient groups need to be incorporated. For example, dibenzo[a,e]-pentalene-embedded thienuquinoids with dicyanomethylene termini were prepared in 2015 by Takimiya’s group.¹¹ They have low-lying LUMO energy levels (−4.2 to −4.3 eV) and showed n-type field-effect transistor (FET) behavior. Under ambient conditions, the devices produced by solution-processing technique showed electron mobility of up to 4.5 × 10⁻² cm² V⁻¹ s⁻¹. Besides this work, to the best of our knowledge, no more n-type pentalene-based materials have been reported.

Generally, to obtain the electron-transporting materials, the electron-deficient imide group is used to lower the energy level and stabilize the radical anions. Therefore, aromatic diimides such as pyromellitic diimide,⁶ naphthalene diimides,⁷ perylene diimides,⁸ longer acene diimides⁹,¹² and large ovalene diimide¹³ were reported to exhibit relatively low energy levels, high electron mobilities, and excellent chemical and thermal stabilities. Following the above design strategies, we expect that incorporation of the imide groups into the diacenopentalene framework would result in new electron-transporting materials with low-lying LUMO energy levels. In this context, two diacenopentalene dicarboximides (DBPDI and DNPD, Scheme 1) were synthesized, and their performance in organic field-effect transistors (OFETs) was evaluated. Both compounds showed n-type FET behavior with moderate electron mobilities.

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DBPDI and DNPDI were measured in chloroform solution. The spectra are shown in Fig. 1a, and the relevant data are summarized in Table 1. Two broad absorption bands located in the visible region were observed for both compounds. The maximum absorption wavelengths of DBPDI and DNPDI in the high-energy region are located at 296 and 328 nm, with molar absorption coefficients of 93,500 and 88,600 M⁻¹ cm⁻¹, respectively. Time-dependent density functional theory (TDDFT) calculations (B3LYP/6-31G*) indicate that this band originates from a combination of multiple HOMO→n → LUMO+m transitions. The bands in the low-energy region with maxima at 485 and 518 nm are observed for DBPDI and DNPDI, respectively, and they mainly come from the HOMO→1 → LUMO transition (DBPDI: λ = 488.2 nm, oscillator strength f = 0.3379) or HOMO → LUMO transition (DNPDI: λ = 512.3 nm, f = 0.5057). The optical energy band gap (E_{Opt}^g) of 2.42 and 2.25 eV can be calculated based on the lowest energy of absorption edge in solution (see Table 1). Compared to DBPDI, the absorption maximum of DNPDI in solution is red-shifted 33 nm due to its more extended π-conjugation.

Electrochemical properties

Cyclic voltammetry (CV) and differential pulse voltammetry (DPV) were used to study the electrochemical properties of compounds DBPDI and DNPDI (Fig. 1b, Table 1 and Fig. S3 in ESI†). The potential was externally calibrated against the ferrocene/ferrocenium couple. Compound DBPDI shows one oxidation wave with half-wave potential (E_{1/2}^{ox}) of 1.36 V and two reversible reduction waves with half-wave potentials E_{1/2}^{red} at −1.46 and −1.11 V. Meanwhile, one oxidation wave with E_{1/2}^{ox} at 1.03 V and two reversible reduction waves with E_{1/2}^{red} at −1.71 and −1.47 V were observed for molecule DNPDI. The HOMO

![Graph showing UV-vis absorption spectra of DBPDI and DNPDI in chloroform solution.](image-url)
and LUMO energy levels were calculated using the following equations: HOMO = [E_{\text{onset}}^{\text{ox}} + 4.8] eV, LUMO = [E_{\text{onset}}^{\text{red}} + 4.8] eV, where E_{\text{onset}}^{\text{ox}} and E_{\text{onset}}^{\text{red}} are the onset of the first oxidation and reduction wave, respectively. The HOMO/LUMO energy levels are calculated to be −5.84/−3.76 eV for DBPDI and −5.72/−3.45 eV for DNPDI. The corresponding electrochemical energy gaps \( E_g^{\text{EC}} \) (LUMO–HOMO) are estimated to be 2.08 eV and 2.27 eV for DBPDI and DNPDI, which are close to their optical band gaps \( E_g^{\text{opt}} \).

**DFT calculations**

DFT calculations were conducted to better understand the electronic properties of DBPDI and DNPDI (Fig. 2 and Fig. S1, S2 and Tables S1, S2 in ESI†). The coefficients for HOMO of DBPDI and HOMO–1 of DNPDI are distributed along the diacenopentane framework, while the phenyl rings are nearly nonconjugated to the backbone. The HOMO–1 of DBPDI and HOMO of DNPDI coefficients are mainly distributed along the zigzag edges of diacenopentene including the phenyl rings. The LUMO coefficient is mainly located on the central pentalene unit and the imide moieties. The HOMO/LUMO energy levels were calculated to be −6.05/−3.31 eV for DBPDI and −5.69/−2.98 eV for DNPDI. It is worth noting that the HOMO and HOMO−1 (−6.15 eV) energy levels of DNPDI are quite close. TDDFT calculations predicted the longest wavelength for HOMO−1 \( \rightarrow \) LUMO transition at 488.2 nm for DBPDI and for HOMO \( \rightarrow \) LUMO transition at 512.3 nm for DNPDI (Fig. S1 and S2 in the ESI†), which is consistent with the experimental data.

**Thermal properties**

Thermal stability is one of the key requirements for the practical application of organic electronic materials. Compounds DBPDI and DNPDI showed good thermal stability, with decomposition temperatures \( T_{\text{db}} \) corresponding to a 5% weight loss in thermogravimetric analysis (TGA) curves at 419 and 460 °C, respectively (Fig. S4a and c, ESI†). The melting points of DBPDI and DNPDI were observed at 354 and 387 °C, and the corresponding crystallization temperatures were observed at 337 and 369 °C, respectively, from differential scanning calorimetry (DSC) curves (Fig. S4b and d, ESI†).

**Organic field effect transistors**

To probe the charge transport properties of compounds DBPDI and DNPDI, we fabricated field-effect transistor (FET) devices for both compounds by solution-processing method. The bottom-gate, top-contact FETs were fabricated on p+Si/SiO\(_2\) substrates by spin-coating 1 wt% chloroform (CHCl\(_3\)) solutions onto 1-octyltri chlorosilane (OTS) or hexamethyldisilazane (HMDS)-treated substrates. The thin films were then annealed at selected temperatures for 10 min in a N\(_2\) glovebox. Au source/drain electrodes (100 nm) were patterned on the organic layer through a shadow mask to produce the devices. The typical transfer and output curves measured in N\(_2\) are shown in Fig. 3. Well-defined saturation characteristics were observed for both. The OFET device parameters for the thin films of DBPDI and DNPDI under different conditions measured under nitrogen are summarized in Table 2. The device parameters are averaged based on over 36 devices in two different batches. The devices showed good uniformity and reproducibility.

Both compounds exhibited n-type FET behaviour. For compound DBPDI, the devices revealed an average electron mobility of 4.6 \( \times 10^{-3} \) cm\(^2\) V\(^{-1}\) s\(^{-1}\) \( (I_{\text{on}}/I_{\text{off}} = 10^{6}) \) and 0.06 cm\(^2\) V\(^{-1}\) s\(^{-1}\) \( (I_{\text{on}}/I_{\text{off}} = 10^{6}) \) for HMDS- and OTS-treated substrates, respectively. The device based on DNPDI displayed an average saturation mobility of 0.011 cm\(^2\) V\(^{-1}\) s\(^{-1}\) \( (I_{\text{on}}/I_{\text{off}} = 10^{6}) \) and 0.034 cm\(^2\) V\(^{-1}\) s\(^{-1}\) \( (I_{\text{on}}/I_{\text{off}} = 10^{6}) \) for HMDS- and OTS-treated substrates, respectively. The thermal annealing has a lower effect on the charge carrier mobilities, indicating that these two materials are good choices for annealing-free device fabrication in the plastic electronics. The DBPDI showed slightly higher charge carrier mobilities and less...
hysteresis, indicating that less traps exist in the thin film due to better film crystallinity and less grain boundaries, which is explained in the following section. Both DBPDI and DNPDI showed decreased charge carrier mobility when their devices were operated in ambient conditions.

Thin-film morphology and solid-state microstructure were characterized by tapping-mode atomic force microscopy (AFM) and 2D X-ray diffraction (XRD). The thin films exhibited ribbon-like crystals for DBPDI and fibre-like crystals for DNPDI (Fig. 4). The root mean square (RMS) values of surface roughness are 29.1 nm and 30.3 nm for DBPDI before and after annealing, respectively. For DNPDI thin film, the RMS values are 7.7 nm and 8.6 nm before and after annealing, respectively. The larger surface roughness of DBPDI thin film is ascribed to the larger crystal size, which reduces the grain boundaries. The XRD measurements of the thin films on OTS-treated substrates exhibited primary peaks at 2θ = 5.8° and 4.5°, which correspond to d-spacing of 15.2 Å and 19.6 Å for DBPDI and DNPDI, respectively (Fig. 5). These are quite close to their molecular lengths of 15.7 Å and 20.6 Å, indicating that both of them are packed in a layer-like structure. In addition, the intensity of the diffraction peak of DBPDI is much stronger than that of DNPDI, although they have a similar film thickness (60 to 70 nm). The better film crystallinity and larger crystal size of DBPDI thin films ensure a lower amount of traps due to the reduced grain boundaries and give efficient charge transport between the source and drain electrodes.

Concluding Remarks

In conclusion, two diacenopentalene diimides, DBPDI and DNPDI, have been successfully synthesized as new π-conjugated organic semiconducting materials for OFETs. They have good solubility in common organic solvents and show high thermal stability, with decomposition temperatures above 400 °C. DBPDI and DNPDI have relatively low-lying LUMO energy levels at −3.76 and −3.45 eV, respectively. Their thin-film transistor test indicated that they are n-type semiconductors. DBPDI and DNPDI devices produced by solution processing technique displayed average electron mobilities of up to 0.06 cm² V⁻¹ s⁻¹ with Ion/Ioff ratio of 2.9 × 10⁶ and 0.034 cm² V⁻¹ s⁻¹ with Ion/Ioff ratio of 1.1 × 10⁵ in N₂, respectively. More effort will be carried out to modify this kind of molecule and optimize fabrication conditions¹⁰ to improve device performance.

Table 2 OFET characteristics of DBPDI- and DNPDI-based devices fabricated on Si/SiO₂ substrates with different surface treatments and different annealing temperatures

| Surface treatment | Annealing temp. | μ [cm² V⁻¹ s⁻¹] | Vg [V] | On/off |
|-------------------|-----------------|----------------|--------|--------|
| DBPDI             | HMDS            | 3.0 × 10⁻³     | 2–4    | 9.6 × 10⁵ |
|                   | XA              | 4.6 × 10⁻³     | 2–4    | 7.8 × 10⁵ |
|                   | OTS-C8          | 0.05           | 7–10   | 2.6 × 10⁶ |
|                   | A120            | 0.06           | 5–8    | 2.9 × 10⁶ |
| DNPDI             | HMDS            | 4.7 × 10⁻³     | 28–30  | 3.2 × 10⁶ |
|                   | XA              | 0.011          | 24–26  | 2.0 × 10⁶ |
|                   | OTS-C8          | 6.8 × 10⁻³     | 23–25  | 2.1 × 10⁵ |
|                   | A160            | 0.025          | 16–18  | 7.8 × 10⁶ |
|                   | A120            | 0.034          | 15–17  | 1.1 × 10⁶ |
Experiment section

Materials

All reagents were purchased from commercial sources without further purification. Anhydrous dichloromethane (DCM) was distilled from CaH₂. 1,4-Dioxane and THF were distilled from sodium-benzophenone immediately prior to use. 4,5-Dibromophthalic acid 1, 1,2-dibromo-4,5-bis(dibromomethyl)benzene 4 and 1-octyl-1H-pyrrole-2,5-dione 5 were prepared by following literature procedures.

General characterization method

¹H and ¹³C NMR spectra were recorded using an Avance 500 MHz Bruker spectrometer in CDCl₃ with tetramethylsilane (TMS) as the internal standard. The chemical shift was recorded in ppm, and the following abbreviations are used to explain the multiplicities: s = singlet, d = doublet, t = triplet, m = multiplet, br = broad. Column chromatography was performed on silica gel 60 (Merck 40–60 nm, 230–400 mesh). EI mass spectra were recorded on an Agilent 5975 C DIP/MS mass spectrometer. UV-vis absorption was recorded on a Shimadzu UV-1700 spectrophotometer. Cyclic voltammetry and differential pulse voltammetry measurements were performed in HPLC-grade dry dichloromethane on a CHI 620C electrochemical analyzer with a three-electrode cell, using 0.1 M Bu₄NPF₆ as the supporting electrolyte, AgCl/Ag as the reference electrode, a gold disk as the working electrode, and Pt wire as the counter electrode, at a scan rate of 100 mV s⁻¹. The potential was externally calibrated against the ferrocene/ferrocenium couple. Differential scanning calorimetry (DSC) was performed on a TA instrument 2920 at a heating rate of 10 °C min⁻¹ under nitrogen flow.

Synthesis

Compound 2 5,6-dibromo-2-hexylisoindoline-1,3-dione. A mixture of 1 (4.0 g, 12.35 mmol) and thionyl chloride (6 mL) in 30 mL of dry dichloromethane was stirred overnight at reflux temperature under argon atmosphere. After removing excess thionyl chloride and solvent, the yellowish solid was directly dissolved in 30 mL of dry acetic acid and used in the next reaction. The hexyl amine (1.3 g, 13.58 mmol) was slowly added, and the mixture was heated to 135 °C overnight under argon. The mixture was cooled to room temperature and extracted with EtAc (80 mL × 2). The combined organic phase was washed with brine (100 mL × 2) and saturated NaHCO₃ solution (100 mL) and dried over anhydrous Na₂SO₄. The organic solvent was removed under reduced pressure, and the residue was purified by column chromatography (silica gel, DCM/hexane = 1 : 1) to afford compound 2 (1.75 g) in 37% yield. ¹H NMR (500 MHz, CDCl₃, ppm) δ = 8.07 (s, 2H), 3.66 (t, J = 7.5 Hz, 2H), 1.68–1.60 (m, 2H), 1.33–1.25 (m, 6H), 0.87 (t, J = 6.8 Hz, 3H), ¹³C NMR (CDCl₃, 125 MHz): δ = 166.47, 131.87, 131.23, 128.33, 38.52, 31.28, 28.37, 26.46, 22.47, 13.95. HR MS (EI): calcd for C₁₄H₁₄Br₂NO₂ (M⁺), 386.9470; found, 386.9485 (error: -0.03 ppm).

Compound 3 5-bromo-2-hexyl-6-(phenylethynyl)isoindoline-1,3-dione. A suspension of Pd(PPh₃)₄Cl₂ (135 mg, 5 mol%) and CuI (36 mg, 5 mol%) and 2 (1.50 g, 3.88 mmol) were purged with argon. After dissolving the mixture with 20 mL of tetrahydrofuran (THF) and 20 mL of anhydrous triethylamine (Et₃N), phenyl acetylene (0.435 g, 4.26 mmol) was slowly added to the solution, then the mixture was stirred at 75 °C overnight. After cooling down to room temperature, the mixture was poured into EtOAc (60 mL × 2) and washed with 10% hydrochloric acid solution (60 mL × 2). The organic layer was dried on Na₂SO₄ and concentrated under vacuum. Finally, the residue was purified carefully by column chromatography (silica gel, DCM/hexane = 1 : 1) to give compound 3 (530 mg) in 34% yield. ¹H NMR (500 MHz, CDCl₃, ppm) δ = 8.07 (s, 1H), 7.97 (s, 1H), 7.63–7.59 (m, 2H), 7.43–7.38 (m, 3H), 3.67 (t, J = 7.3 Hz, 2H), 1.70–1.62 (m, 2H), 1.34–1.25 (m, 6H), 0.88 (t, J = 7.0 Hz, 3H), ¹³C NMR (CDCl₃, 125 MHz): δ = 167.15, 166.69, 131.96, 131.63, 131.58, 131.07, 130.82, 129.57, 128.56, 127.34, 127.26, 121.92, 98.69, 87.20, 34.84, 31.31, 28.44, 26.49, 22.48, 13.95. HR MS (EI): calcd for C₁₄H₁₂BrNO₂ (M⁺), 409.0667; found, 409.0669 (error: -0.16 ppm).

Compound DBPDI. A reaction flask was charged with hydroquinone (0.113 g, 1.20 mmol), Cs₂CO₃ (0.334 g, 1.02 mmol), CsF (0.171 g, 1.13 mmol), P(2-furyl)₃ (15 mg, 0.08 mmol), and Pd₃dba₃ (15 mg, 0.02 mmol) and purged with argon. A solution of 3 (0.200 g, 0.50 mmol) in 1,4-dioxane (15 mL) was injected into the catalyst mixture with a syringe. The suspension was immediately heated to 135 °C. After heating for 24 h, the reaction mixture was diluted with CH₂Cl₂ (20 mL), filtered through celite, and then concentrated to yield raw solid. After purification via column chromatography (silica gel, DCM/hexane = 1 : 4), the product was dissolved in CHCl₃, precipitated in methanol/acetone = 3 : 2 and filtered. This procedure was repeated three times to give the purer product DBPDI (12 mg) in 10% yield for characterization. ¹H NMR (500 MHz, CDCl₃, ppm) δ = 7.65–7.57 (m, 10H), 7.56 (s, 2H), 7.49 (s, 2H), 3.61 (t, J = 7.3 Hz, 4H), 1.66–1.58 (m, 4H), 1.29–1.25 (m, 12H), 0.86 (t, J = 6.8 Hz, 6H), ¹³C NMR (CDCl₃, 125 MHz): δ = 168.17, 168.05, 154.70, 143.98, 143.44, 140.43, 132.51, 132.45, 131.77, 130.42, 129.47, 128.31, 117.70, 116.41, 38.11, 31.34, 28.50, 26.46, 22.48, 13.98. HR MS (EI): calcd for C₁₄H₁₄BrNO₂ (M⁺), 660.2988; found, 660.2994 (error: 0.91 ppm).

Compound 6,7-dibromo-2-octyl-1H-benzo[f]isoindole-1,3(2H)-dione. NaI (1.55 g, 10.5 mmol) was added with stirring to a solution of 4 (1.20 g, 2.1 mmol) and N-octylmaleimide 5 (0.50 g, 2.39 mmol) in N,N-dimethylethamide (20 mL). The mixture was heated at 80 °C for 12 h. The product was collected from the dark solution by filtration and washed with water, methanol and acetone. The residue was purified by flash column chromatography (silica gel, DCM/hexane = 1 : 1) to provide compound 6 (0.85 g) in 88% yield. ¹H NMR (500 MHz, CDCl₃, ppm) δ = 8.33 (s, 2H), 8.20 (s, 2H), 3.74 (t, J = 7.5 Hz, 2H), 1.73–1.68 (m, 2H), 1.37–1.20 (m, 10H), 0.86 (t, J = 7.0 Hz, 3H), ¹³C NMR (CDCl₃, 125 MHz): δ = 167.36, 134.78, 134.18, 129.12, 126.15, 123.27, 38.55, 31.75, 29.13, 28.47, 26.90, 22.60, 14.04. HR MS (EI): calcd for C₂₉H₂₃Br₂NO₂ (M⁺), 464.9399; found, 464.9391 (error: 2.58 ppm).

Compound 7 6-bromo-2-octyl-7-(phenylethynyl)-1H-benzo[f]isoindole-1,3(2H)-dione. A suspension of Pd[PPh₃]₄Cl₂ (65 mg, 5 mol%), CuI (15 mg, 5 mol%), and 6 (850 mg, 1.8 mmol) was purged with argon. Dissolving the mixture with 15 mL of
anhydrous tetrahydrofuran (THF) and 15 mL anhydrous triethylamine (Et3N), phenyl acetylene (180 mg, 1.8 mmol) was slowly added to the solution, then the mixture was stirred at 75 °C overnight. After cooling down to room temperature, the mixture was poured into EtOAc (100 mL) and extracted with diluted hydrochloric acid (60 mL × 3). The organic layer was dried on Na2SO4 and concentrated under vacuum. Finally, the residue was purified carefully by column chromatography (silica gel, DCM/hexane = 1:10) to give compound 7 (380 mg) in 43% yield. 1H NMR (500 MHz, CDCl3, ppm) δ = 8.32 (s, 1H), 8.26 (s, 1H), 8.24 (s, 1H), 8.21 (s, 1H), 7.66–7.62 (m, 2H), 7.43–7.40 (m, 3H), 3.74 (t, J = 7.5 Hz, 2H), 1.74–1.67 (m, 2H), 1.40–1.20 (m, 10H), 0.87 (t, J = 6.8 Hz, 3H). 13C NMR (CDCl3, 125 MHz): δ = 167.58, 167.49, 135.25, 134.28, 133.79, 133.16, 134.88, 129.24, 129.22, 128.82, 128.52, 126.37, 125.95, 123.94, 123.19, 122.30, 96.38, 87.42, 81.30, 71.26, 69.51. HR MS (EI): calcd for C30H28BrNO2 (M+), 503.1117; found, 503.1116 (error: +0.19 ppm).

**Compound DNPDI.** A reaction flask was charged with hydroquinone (0.172 g, 1.83 mmol), Cs2CO3 (0.508 g, 1.55 mmol), CsF (0.260 g, 1.72 mmol), P(2-furyl)3 (23 mg, 0.12 mmol), and Pd2(dba)3 (23 mg, 0.026 mmol) and purged with argon. A solution of the catalyst mixture with a syringe. The suspension was immediately heated to 135 °C. After heating for 24 h, the reaction mixture was diluted with CH2Cl2 (25 mL), filtered through celite, and then concentrated under vacuum to yield red solid. After purification via column chromatography (silica gel, DCM/hexane = 1:4), the product was dissolved in CHCl3, precipitated in methanol/acetone (3:2) and filtered. This procedure was repeated three times to give the purer product **DNPDI** (75 mg) in 24% yield for characterization. 1H NMR (500 MHz, CDCl3, ppm) δ = 8.08 (s, 2H), 8.05 (s, 2H), 7.98 (s, 2H), 7.83 (d, J = 7.0 Hz, 4H), 7.73 (s, 2H), 7.69 (t, J = 7.5 Hz, 4H), 7.66–7.61 (m, 2H), 3.67 (t, J = 7.0 Hz, 4H), 1.68 (br, 4H), 1.53–1.20 (m, 20H), 0.86 (t, J = 6.5 Hz, 6H). 13C NMR (CDCl3, 125 MHz): δ = 167.85, 150.29, 144.83, 139.78, 136.67, 136.01, 134.77, 133.19, 129.80, 129.34, 129.03, 128.77, 128.72, 128.46, 124.70, 123.45, 123.33, 38.41, 31.80, 29.17, 29.15, 28.58, 27.00, 22.59, 13.93. HR MS (EI): calcd for C56H52N2O4 (M+), 875.3927; found, 875.3926 (error: −0.23 ppm).

**Fabrication and characterization of OFET devices**

Top-contact, bottom-gate TFTs were prepared. A heavily p++ doped silicon wafer (100, Silicon Quest International, resistivity <0.005 Ω cm−1) with 200 nm thermal silicon dioxide (SiO2) was used as the substrate/gate electrode, with the SiO2 layer serving as the gate dielectric. The SiO2/Si substrate was cleaned with acetone and isopropanol (IPA). It was then immersed in a piranha solution (V(H2SO4):V(H2O2) = 2:1) for 20 minutes, followed by rinsing with deionized water, and then re-immersed in 0.1 M OTS or HMDS in anhydrous toluene at 60 °C for 20 minutes. It was then rinsed with toluene, and then blow-dried with nitrogen gas. The semiconductor layer was deposited on top of the OTS- or HMDS-modified substrates by spin-coating the 1.0 wt% solution in chloroform, then thermal annealing at selected temperatures for 20 min. Subsequently, gold source/drain electrode pairs were deposited by thermal evaporation through a metal shadow mask to create a series of OFETs with various channel length (L = 100/150 mm) and width (W = 1 mm) dimensions. The OFET devices were then characterized using a Keithley SCS-4200 probe station under N2 conditions in the dark.

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