Linear-type carbazoledioxazine-based organic semiconductors: the effect of backbone planarity on the molecular orientation and charge transport properties†

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We report the synthesis of a linear-type dibromocarbazoledioxazine (CZ) derivative as a new precursor for semiconducting polymers. The chemical structures of the CZ unit and its polymers with thiophene or thienothiophene spacers (namely, PCZT and PCZTT) were fully characterized. PCZT and PCZTT possessed similar medium optical band gap ($E_{\text{opt}}^{\text{final}}$) and electrochemical band gap ($E_{\text{opt}}^{\text{CV}}$) of around 1.70 eV estimated from the onset absorption and electrochemical redox potentials of the thin films, respectively. Computational density functional theory (DFT) calculations suggested that the backbone of the PCZT might be highly twisted, while that of PCZTT could be very planar. The effect of different backbone geometries on the charge-transport properties was studied by using thin film transistors (TFTs). The TFT device based on PCZTT showed a four times higher hole mobility as compared to that based on PCZT. The superior TFT performances of PCZTT were reasonably attributed to its edge-on backbone packing orientations toward the Si substrate revealed by the grazing-incidence wide-angle X-ray scattering (GIWAXS), which was favorable for in-plane charge transport in the TFT devices.

Among the reported acceptor building blocks, dye/pigment molecules have received enormous interest. Diketopyrrolopyrrole (DPP), isoindigo (IID), and rylene diimide (such as naphthalenediimide (NDI) and perylenediimide (PDI)) are successful examples for the construction of high-performance semiconducting polymers with mobilities ($\mu$) > 1 cm$^2$ V$^{-1}$ s$^{-1}$ in organic TFTs and power conversion efficiencies (PCE) > 10% in OPVs.

Among these dye/pigment molecules, the angular-type carbazoledioxazine (CZ, X = Cl, Chart 1 middle), also called pigment violet 23, is a well-known commercial pigment, but it has been less explored in the field of organic electronics. Note that this pigment is not only a member of the electron-accepting dioxazine family, but also originates from the electron-donating carbazole. Thus, the violet color is derived from the highly-efficient intramolecular charge-transfer (CT) band. The general synthesis of this pigment is based on the condensation of a 3-aminocarbazole derivative with $p$-chloranil followed by two-fold cyclization reactions. In the final cyclization process, simple heating selectively affords the angular-type CZ (Chart 1 middle). On the other hand, the introduction of the methoxy-substituent into the 2-position of the carbazole unit produces the linear-type CZ under the same conditions (Chart 1 middle). The electrochemical and optical properties of some CZ derivatives were investigated in the 1990s. However, there have been, to the best of our knowledge, no reports about their applications in organic electronic devices until our recent...
producing two promising semiconductors, namely ditin

Experimental

General measurements

Nuclear magnetic resonance (NMR) spectra were recorded using a JEOL model AL300 (300 MHz) at room temperature. Deuterated chloroform was used as the solvent. Chemical shifts of NMR were reported in ppm (parts per million) relative to the residual solvent peak at 7.26 ppm for $^1$H NMR spectroscopy and 77.6 ppm for $^{13}$C NMR spectroscopy. Coupling constants ($J$) were given in Hz. The resonance multiplicity was described as $s$ (singlet), $d$ (doublet), $t$ (triplet), and $m$ (multiplet). Fourier transform infrared (FT-IR) spectra were recorded on a JASCO FT/IR-4100 spectrometer in the range from 4000 to 600 cm$^{-1}$. MALDI-TOF mass spectra of small molecules were measured on a Shimadzu/Kratos AXIMACFR mass spectrometer equipped with a nitrogen laser ($\lambda = 337$ nm) and pulsed ion extraction, which was operated in a linear-positive ion mode at an accelerating potential of 20 kV. Tetrahydrofuran (THF) solutions containing 1 g L$^{-1}$ of a sample, 20 g L$^{-1}$ of dithanol, and 1 g L$^{-1}$ of sodium trifluoroacetate were mixed at a ratio of 1 : 1 ; 1; and then 1 mL aliquot of this mixture was deposited onto a sample target plate. MALDI-TOF mass spectra of polymers were measured on a Bruker UltraflexXtreme mass spectrometer, which was operated in a linear-positive ion mode. Dithanol was used as a matrix. Elemental analysis was conducted at the Center for Advanced Materials Analysis, Tokyo Institute of Technology. The elements of C, H, and N were measured on a J-Science JM10, the elements of Cl, Br, and S on a Yanako HSU-20+ICS-1100, and the element of O on an Elementar Vario micro cube. UV-vis-NIR spectra were recorded on a JASCO V-670 spectrophotometer. Thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC) measurements were carried out on a Rigaku TG8120 and a Rigaku DSC8230, respectively, under nitrogen flow at the scan rate of 10 °C min$^{-1}$. Electrochemistry measurements were carried out on a BAS electrochemical analyzer model 612C at 20 °C in a classical three-electrode cell. The working, reference, and auxiliary electrodes were a glassy carbon electrode, Ag/AgNO$_3$/CH$_3$COONa/nC$_4$H$_9$NPF$_6$, and a Pt wire, respectively. For calibration, the redox potential of ferrocene/ferrocinium (Fc/Fc') was measured. It was assumed that the redox potential of Fe/Fc' has an absolute energy level of $-4.80$ eV to vacuum. The HOMO and LUMO
Energy levels were then calculated according to the following equations:

\[ E_{\text{HOMO}} = -(\varphi_{\text{ox}} + \varphi_{\text{Fc/Fc}^+} + 4.80) \text{ (eV)} \] (1)

\[ E_{\text{LUMO}} = -(\varphi_{\text{re}} + \varphi_{\text{Fc/Fc}^+} + 4.80) \text{ (eV)} \] (2)

where \( \varphi_{\text{ox}} \) is the onset oxidation potential vs. Ag/AgNO\(_3\), \( \varphi_{\text{re}} \) is the onset reduction potential vs. Ag/AgNO\(_3\), and \( \varphi_{\text{Fc/Fc}^+} \) is the redox potential of ferrocene/ferrocenium vs. Ag/AgNO\(_3\) and it is \(-0.05\) V in this study.

**Fabrication and characterization of thin film transistors**

Top-contact/bottom-gate TFT devices were fabricated using n\(^+\)-Si/SiO\(_2\) substrates where n\(^+\)-Si and SiO\(_2\) were used as the gate electrode and gate dielectric, respectively. The substrates were subjected to cleaning with deionized water, acetone, and 2-propanol. Then, the substrates were cleaned with O\(_3\) for 20 min using a Technovision model 208 UV-O\(_3\) cleaning system. The cleaned substrates were modified with octadecyltrimethoxysilane (OTMS) to form a self-assembled monolayer (SAM). Thin films of the CZ-based semiconductors were deposited on the treated substrate by spin-coating a chloroform solution (5 g L\(^{-1}\)), followed by thermal annealing at 150, 200, 250, or 300 °C for 10 min in an argon-filled glove box. After the thin film deposition, ~50 nm thick gold was deposited as source and drain contacts using a shadow mask. The TFT devices had a channel length (\( L \)) of 100 \( \mu \)m and a channel width (\( W \)) of 1 mm. The TFT performances were measured under ambient conditions using a Keithley 4200 parameter analyzer on a probe stage. The carrier mobilities, \( \mu \), were calculated from the data in the saturated regime according to the following equation:

\[ I_{\text{SD}} = \left( \frac{W}{2L} \right) C_{\text{i}} \mu (V_{\text{GS}} - V_{\text{th}})^2 \] (3)

where \( I_{\text{SD}} \) is the drain current in the saturated regime, \( W \) and \( L \) are the semiconductor channel width and length, respectively, \( C_{\text{i}} (C_{\text{i}} = 13.3 \text{ nF cm}^{-2}) \) is the capacitance per unit area of the gate dielectric layer, and \( V_{\text{GS}} \) and \( V_{\text{th}} \) are the gate voltage and threshold voltage, respectively. \( V_{\text{GS}} - V_{\text{th}} \) of the devices was determined from the square root values of \( I_{\text{SD}} \) at the saturated regime. Current on/off ratios (\( I_{\text{on}}/I_{\text{off}} \)) were determined from the minimum current at around \( V_{\text{GS}} = +20 \) to \(-20\) V (\( I_{\text{on}} \)) and the current at \( V_{\text{GS}} = -80 \) V (\( I_{\text{off}} \)). The transfer characteristics were obtained by sweeping the \( V_{\text{GS}} \) from +20 to –80 V.
Grazing-incidence wide angle X-ray scattering (GIWAXS) measurements

The GIWAXS profiles were obtained by using a Phillips X’Pert-Pro-MRD with monochromated CuKα radiation (wavelength = 0.154 nm) operated at 45 kV and 40 mA in air. The samples were exposed to the X-ray beam for 1 h with an incident beam radius of 320 mm. The thin films for GIWAXS were prepared by spin-coating the polymer solutions on OTMS-treated Si/SiO2 substrates followed by thermal annealing at the optimized temperature for 10 min under vacuum.

Atomic force microscopy (AFM) measurements

AFM samples were prepared by spin-coating the polymer solutions on OTMS-treated Si/SiO2 substrates. Both pristine and thermally-treated films were examined by a Seiko Instruments SPA-400 with a stiff cantilever of Seiko Instruments DF-20.

Materials and synthetic details

All chemicals were purchased from Tokyo Chemical Industry (TCI), Kanto Chemical, and Sigma Aldrich and used as received unless otherwise stated. 2-Bromo-7-methoxy-9-carbazole (7.83 g, 20.8 mmol) was added and the mixture was quenched with H2O and extracted with CH2Cl2. A mixture of fuming nitric acid (2.0 mL, 48.2 mmol) and acetic anhydride (80 mL) was slowly added to 2-bromo-7-methoxy-9-carbazole (5.34 g, 8.71 mmol) in acetic acid (80 mL), sodium acetate (0.542 g, 6.61 mmol), and ethanol (2922, 2853, 1738, 1714, 1574, 1363, 1219, 755, 745 cm⁻¹) in dry DMF under nitrogen for 5 h. After stirring for 30 min under nitrogen. A solution of 2,5-bis((7-bromo-9-(2-decyltetradecyl)-2-methoxy-3-nitro-9H-carbazol-3-yl)amino)-3,6-dichlorocyclohexa-2,5-diene (4) Iron dust (22.5 g, 0.403 mol) was added to water (30 mL) and aqueous HCl (10 N, 5 mL), and the mixture was stirred at 90 °C for 30 min under nitrogen. After a solution of 3 (8.64 g, 13.1 mmol) in 2-propanol (150 mL) was added, the mixture was refluxed for 5 h. After cooling to room temperature, water was added and the organic phase was extracted with CH2Cl2. Evaporation yielded 7-bromo-9-(2-decyltetradecyl)-2-methoxy-9H-carbazol-3-amine, and this compound was subjected to the next reaction without further purification. p-Chloranil (1.64 g, 6.67 mmol), sodium acetate (0.542 g, 6.61 mmol), and ethanol (200 mL) were added, and the mixture was refluxed under nitrogen for 5 h. After cooling to room temperature, water was added and the organic phase was extracted with CH2Cl2. Evaporation followed by column chromatography (SiO2, hexane/CH2Cl2 5 : 4) afforded the desired compound as a reddish-purple solid (3.95 g, two-step yield of 42%).

Synthesis of 2,5-bis((7-bromo-9-(2-decyltetradecyl)-2-methoxy-9H-carbazol-3-yl)amino)-3,6-dichlorocyclohexa-2,5-diene (4)
Results and discussion

Synthesis and characterization

Starting from the commercially-available 1,4-dibromo-2-nitrobenzene, 2-bromo-7-methoxy-9H-carbazole (1) was obtained in the moderate yield of 42% in two steps according to a literature report.\(^{46}\) Alkylation at the 9-position followed by mono-nitration yielded 7-bromo-9-(2-decyttetradecyl)-2-methoxy-3-nitro-9H-carbazole (3) in high yield (two steps, total yield over 70%, Scheme 1). The reduction of 3 by hydrochloric acid and Fe powder afforded the corresponding 3-amino-carbazole derivative, and this compound was directly used for the next condensation reaction with p-chloranil (2,3,5,6-tetrachlorocyclohexa-2,5-diene-1,4-dione) due to the limited stability of amine compound in air. The resulting product (4) was further subjected to a ring closure reaction at high temperature (220 °C) in 1-chloronaphthalene as the solvent, leading to the target linear carbazoledioxazine monomer (CZ) in 28% isolated yield after multiple purifications by Soxhlet extraction and recycling preparative high performance liquid chromatography (HPLC). The chemical structure of the new bifunctional monomer was unambiguously characterized by \(^{1}\text{H}, \text{C}\) NMR, IR spectroscopies and MALDI-TOF mass spectrometry. Particular attention was paid to the isomeric structures of this product. The resonance multiplicity of a set of aromatic proton peaks was observed as a singlet (protons d and g in Fig. 1 or protons a and b in Fig. S1†), clearly suggesting the formation of the linear-type structure. This was also consistent with the reported spectral patterns of similar derivatives.\(^{35-48}\) In addition, after ring closure, the proton peaks of N–H and OCH\(_3\) as well as the IR peaks ascribed to the C=O and NH vibrations completely disappeared (Fig. 1 and S2†).

The linear-type dibromocarbazole-dioxazine monomer (CZ) was then reacted with the ditin compounds of thiophene (T) and thienothiophene (TT) by the conventional Stille polycondensation to afford the target products, namely PCZT and PCZTT, respectively (Scheme 1). They were washed using a Soxhlet extractor with the sequence of methanol, hexane, and acetone to remove the low molecular weight oligomers. Finally, the chloroform soluble fractions were collected and were found to be reasonably soluble in a variety of chlorinated solvents, such as chloroform, chlorobenzene, and 1,2-dichlorobenzene at room temperature. Thus, the molecular weights (number-average molecular weight (M\(_n\)) and weight-average molecular weight (M\(_w\))) were determined by gel permeation chromatography (GPC) measurements using polystyrene as the standard and 1,2-dichlorobenzene as the eluent at 40 °C. Both polymers showed similar molecular weights and polydispersity indices.
(PDI, $M_w/M_n$): PCZT ($M_n = 6.2$ kg mol$^{-1}$, $M_w = 15.5$ kg mol$^{-1}$, PDI = 2.5) and PCZTT ($M_n = 6.8$ kg mol$^{-1}$, $M_w = 15.4$ kg mol$^{-1}$, PDI = 2.3). The $M_n$ values of PCZT and PCZTT suggested the pentamers (Table 1 and Fig. S3†). This was also supported by the comprehensive elemental analysis of both polymers. Importantly, a small amount of Br was detected (1.74% for PCZT and 2.87% for PCZTT). Under the assumption that the both terminal units are Br-CZ, the $M_n$ was calculated to be 9.1 kg mol$^{-1}$ for PCZT and 5.5 kg mol$^{-1}$ for PCZTT, which were consistent with the GPC results. These results also suggested that the rate-determining step of the Stille polycondensation is oxidative character verification. As shown in Fig. 2, one could observe some sets of molecular ion peaks ascribed to the desired repeat unit structures. These spectra support the progress of the Stille polycondensation at the Br moieties of the CZ monomer and rule out the possibility of undesired side reactions at the Cl moieties. It should be noted that the peak intensities of the MALDI-TOF mass spectra significantly decrease as the molecular weights increase, although they are dependent on the sample purity, molecular weight distributions, and matrix and instrument settings.

**Thermal properties**

The thermal stability of the polymers was estimated by thermogravimetric analysis (TGA) under a nitrogen atmosphere at the heating rate of 10 °C min$^{-1}$ (Fig. S4†). Both PCZT and PCZTT exhibited a similar thermal stability profile with the 5% weight loss temperatures ($T_{5\%}$) being ~365 °C (365 °C for PCZT and 368 °C for PCZTT). The $T_{d}$ values were sufficiently high for advanced studies of the TFT fabrication, which requires optimization of the thermal annealing conditions. The thermal transition properties were also investigated by differential scanning calorimetry (DSC) measurements in the temperature range without thermal decomposition under N$_2$ at the heating or cooling rate of 10 °C min$^{-1}$ (Fig. S5†). Although no sharp transitions appeared, both polymers showed some endothermic

### Table 1 Molecular weights, optical and electrochemical properties of the CZ monomer and its polymers

<table>
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<tr>
<th></th>
<th>$M_n$ $^a$ (g mol$^{-1}$)</th>
<th>PDI $^a$</th>
<th>DP $^b$</th>
<th>$\lambda_{\text{sol}}$ (nm)</th>
<th>$\lambda_{\text{film}}$ $^b$ (nm)</th>
<th>$\lambda_{\text{onset}}$ (nm)</th>
<th>$\varepsilon_{\text{opt}}$ $^c$ (eV)</th>
<th>$\varphi_{\text{on}}$ $^d$ (V)</th>
<th>$\varphi_{\text{on}}$ $^d$ (eV)</th>
<th>$E_{\text{HOMO}}$ (eV)</th>
<th>$E_{\text{LUMO}}$ (eV)</th>
<th>$\Delta E_{\text{opt}}$ $^e$ (eV)</th>
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<td>—</td>
<td>554, 599</td>
<td>573, 629</td>
<td>702</td>
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<td>-5.27</td>
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<td>6200</td>
<td>2.5</td>
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<td>389, 620</td>
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<td>0.42</td>
<td>-1.28</td>
<td>-5.17</td>
<td>-3.47</td>
<td>1.70</td>
</tr>
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<td>PCZTT</td>
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<td>2.3</td>
<td>5</td>
<td>412, 624</td>
<td>433, 677</td>
<td>770</td>
<td>1.61</td>
<td>0.35</td>
<td>-1.29</td>
<td>-5.10</td>
<td>-3.46</td>
<td>1.64</td>
</tr>
</tbody>
</table>

$^a$ Determined by GPC. $^b$ Cast onto the glass substrate. $^c$ Optical band gap ($E_{\text{opt}}$) estimated from the onset wavelength of the films. $^d$ Thin films measured in CH$_3$CN with 0.1 M (nC$_4$H$_{10}$)$_2$NCIO$_4$ at the scan rate of 0.1 V s$^{-1}$. Potentials vs. Fe/Fe$^+$. $^e$ Theoretical.

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**Fig. 2** MALDI-TOF mass spectra of (a) PCZT and (b) PCZTT.
and exothermic peaks during the heating and cooling processes, respectively. For example, PCZT displayed a transition at 208 °C in the second heating process, suggesting the existence of some different phases. PCZTT also showed a similar phase behavior.14,15

**Optical and electrochemical properties**

The absorption spectra of CZ, PCZT, and PCZTT in both solutions and thin films are shown in Fig. 3a. CZ displayed two absorption peaks with an optical band gap (E\(_{\text{opt}}\)) of 1.77 eV estimated from the onset wavelength (\(\lambda_{\text{onset}}\)) of the film (Table 1). After reacting with the thiophene or thienothiophene co-monomer units, PCZT and PCZTT further extended the effective conjugation length, leading to red-shifted absorption with a narrower band gap of 1.67 and 1.61 eV, respectively. The absorption maxima in dilute 1,2-dichlorobenzene solution and in the thin film state (\(\lambda_{\text{sol}}\) and \(\lambda_{\text{film}}\)) of PCZTT bathochromically shifted as compared to those of PCZT. For example, the film absorption (\(\lambda_{\text{film}}\)) of PCZTT was detected at 433 and 677 nm, while the PCZT film showed the \(\lambda_{\text{film}}\) of 399 and 663 nm (Table 1). Accordingly, PCZTT possessed a longer \(\lambda_{\text{onset}}\) and narrower \(E_g\) than PCZT. This superior optical property of PCZTT was reasonably attributed to the stronger electron-donating nature of the thienothiophene unit, which elongates the effective conjugation length and raises the HOMO level.4,13

To determine the frontier energy levels, cyclic voltamograms (CVs) were measured as shown in Fig. 3b and the corresponding data are listed in Table 1. Thin films of the samples were prepared on a glassy carbon electrode, and their CVs were measured in CH\(_3\)CN with 0.1 M \((n\text{C}_4\text{H}_9)_4\text{NClO}_4\) at room temperature under an argon atmosphere.12 All three samples displayed a well-defined reversible oxidation peak and an irreversible reduction one. The oxidation currents were much higher than the corresponding reduction ones (Fig. 3b), indicating the more facilitated hole injection to these thin films.11,12 The highest occupied molecular orbital (E\(_{\text{HOMO}}\)) and lowest unoccupied molecular orbital (E\(_{\text{LUMO}}\)) values were estimated from the onset oxidation (\(\varphi_{\text{ox}}\)) and onset reduction (\(\varphi_{\text{re}}\)) potentials, respectively. As listed in Table 1, the monomer CZ possessed the E\(_{\text{HOMO}}/E_{\text{LUMO}}\) of \(-5.27/-3.50 \text{ eV}\), which were comparable to its angular type counterparts.49 The thin film of PCZT showed the \(\varphi_{\text{ox}}\) of 0.42 V and \(\varphi_{\text{re}}\) of \(-1.28 \text{ V}\), corresponding to the E\(_{\text{HOMO}}\) value of \(-5.17 \text{ eV}\) and the E\(_{\text{LUMO}}\) value of \(-3.47 \text{ eV}\). The frontier energy levels of PCZTT were similarly determined. While the HOMO level of PCZTT was 0.07 eV higher than that of PCZT, the LUMO levels of both polymers were almost the same within experimental error (merely 0.01 eV). The electrochemical band gaps (E\(_g\)) of both polymers, calculated from the \(\varphi_{\text{ox}}\) and \(\varphi_{\text{re}}\) values, correlated with their E\(_{\text{opt}}\) (Table 1). For example, the E\(_g\) values of both polymers were 0.03 eV higher than the corresponding E\(_{\text{opt}}\).11

In order to further understand the optoelectronic properties and molecular geometry of the new polymers, density functional theory (DFT) calculations were performed using the

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**Fig. 3**  (a) UV-vis-NIR absorption profiles of CZ, PCZT, and PCZTT in dilute 1,2-dichlorobenzene solution and thin film states spin-cast on a glass substrate; (b) cyclic voltammograms of the CZ, PCZT, and PCZTT films drop-cast on a glassy carbon electrode, measured in acetonitrile with 0.1 M \((n\text{C}_4\text{H}_9)_4\text{NClO}_4\) at the scan rate of 0.1 V s\(^{-1}\); (c) calculated HOMO/LUMO orbitals and side-view geometry of PCZT; and (d) calculated HOMO/LUMO orbitals and side-view geometry of PCZTT (using DFT B3LYP/6-31G(d), long and branched alkyl chains are substituted by the methyl group).
dimer structures as the models with a B3LYP/6-31G(d) basis set. The N-alkyl substituents and alkyl side chains were simplified as a methyl group. The molecular orbitals (MOs) and side-view geometry are shown in Fig. 3c and d. One could observe that the HOMO orbitals were well distributed along the backbone, while the LUMO orbitals were located on one side (mainly on the acceptor units). This kind of HOMO/LUMO distribution well explains the CV results. The radical cations formed through the one-electron oxidation are stable due to the localized LUMO on the backbone, while the LUMO orbitals were located on one side (mainly on the acceptor units). This kind of HOMO/LUMO distribution well explains the CV results. The radical cations formed through the one-electron oxidation are stable due to the localized LUMO on the acceptor unit. Interestingly, there was an explicit difference in the optimized backbone geometry between PCZT and PCZTT. The PCZT dimer formed an angular-shape causing a twisted backbone with the dihedral angle of 40°. In contrast, the PCZTT dimer possessed an almost planar backbone due to the negligible steric repulsion between the adjacent repeat units (Fig. 3c and d bottom). These calculations are consistent with the experimental optical and electrochemical results.

Fabrication and measurements of thin film transistors

To further study the different backbone geometries effect on the charge transport properties, organic TFTs with bottom-gate/top-contact structures were fabricated by spin-coating the polymer solutions onto an octadecyltrimethoxysilane (OTMS)-treated SiO₂ (300 nm)/n⁺-Si substrate. After thermal annealing at some selected temperatures (100, 150, 250, or 300 °C), ~50 nm thick gold was deposited as the source and drain contacts using a shadow mask. As anticipated from the determined shallow HOMO (ca. −5.1 eV) and LUMO (ca. −3.5 eV) levels, both polymers showed unipolar p-type transistor performances (Fig. 4a to d). For example, the as-cast films of PCZT and PCZTT exhibited hole mobilities ($\mu_h$) of $1.4 \times 10^{-4}$ and $5.1 \times 10^{-5}$ cm² V⁻¹ s⁻¹, respectively (Fig. 4a and c and Table 2). Subsequently, we tried thermal annealing to optimize the TFT performances. Unexpectedly, the mobility of PCZT decreased to $3.5 \times 10^{-7}$ cm² V⁻¹ s⁻¹ after the thermal annealing at 100–150 °C, but it recovered to $1.4 \times 10^{-4}$ cm² V⁻¹ s⁻¹ when the thermal annealing temperature was raised to 250 °C (Fig. 4e and Table 2). The highest $\mu_h$ of $3.0 \times 10^{-4}$ cm² V⁻¹ s⁻¹ was achieved after thermal annealing at 300 °C (Fig. 4b and e and Table 2). On the other hand, the mobility of PCZTT had a linear correlation with the thermal annealing temperature (Fig. 4e). The highest $\mu_h$ of $1.4 \times 10^{-3}$ cm² V⁻¹ s⁻¹ was achieved with the annealing temperature of 300 °C (Fig. 4d). It should be noted that this value is relatively good, considering the amorphous thin film structures of these polymers (for the GIWAXS results, vide infra). Overall, PCZTT with a planar backbone demonstrated four times higher mobility than that of PCZT with a twisted backbone (Table 2). Further improvements of the mobilities would be possible by the combination with other prominent building blocks to enhance the intramolecular CT character and thin film crystallinity.

Fig. 4 Transfer characteristics of TFTs based on the thin films of (a) PCZT without annealing; (b) PCZT annealed at the optimized temperature of 300 °C; (c) PCZTT without annealing; (d) PCZTT annealed at the optimized temperature of 300 °C ($L = 100 \mu m$ and $W = 1 \ mm$. All the measurements were done under ambient conditions). (e) Effects of annealing temperatures on the hole mobilities of PCZT and PCZTT.
Furthermore, the thermal annealing treatment at 300 °C adopted an edge-on backbone packing orientation (Fig. S8b) with a chain-to-chain packing distance of 18.0 Å (out-of-plane) was found in its 1D pro-
tection peak (100) located at 2θ = 4.0° along the qy direction (in-plane direction). In addition, a reflection peak (010) located at 2θ = 25.7° along the qx direction (out-of-plane) was found in its 1D profile (Fig. S7b†). This result indicated that PCZT tends to adopt a face-on backbone packing orientation with the chain-to-chain and π-π stacking distances of 22.0 Å and 3.5 Å, respectively (Fig. S8a†).13,25 On the other hand, the thin film of PCZTT exhibited a primary diffraction peak [100] located at 2θ = 4.9° along the qz direction (out-of-plane) with a chain-to-chain packing distance of 18.0 Å (Fig. 5c and S7c†). This result suggested that PCZTT tends to adopt an edge-on backbone packing orientation (Fig. S8b†). Furthermore, the thermal annealing treatment at 300 °C explicitly intensified the primary diffraction peak (Fig. 5d and S7d†). Therefore, the higher mobility of PCZTT was reasonably attributed to its edge-on backbone packing orientations toward the Si substrate, which was favorable for the in-plane charge transport in the TFT devices (Fig. S8b†).10,13,14,22,25 Overall, the GIWAXS results were well correlated with the TFT performances, and it was revealed that the polymers with a planar backbone show a better charge carrier transport property than those with a twist backbone.10,13,13,22,25

To elucidate the thermal annealing effects on the thin film quality, the surface morphology of the thin films was investigated using tapping-mode atomic force microscopy (AFM).13,14 As shown in Fig. 6, the thin films of PCZT showed a relatively amorphous morphology. A root-mean-square (RMS) roughness of 0.63 nm was observed for the pristine film (Fig. 6a). Thermal annealing at 150 °C increased the RMS value to 0.86 nm, whereas it decreased to 0.50 nm at the thermal annealing temperature of 300 °C (Fig. 6b and c). A smoother surface morphology would result in a better interfacial contact with the Au electrode, and accordingly, the charge carrier transport would be facilitated.14,22,25 Therefore, the thin film annealed at 150 °C showed the lowest mobility, while annealing at 300 °C led to the highest mobility in the case of PCZT. On the other hand, the RMS values of the PCZTT linearly decreased as the thermal annealing temperature increased. For instance, the pristine film showed a relatively rough surface morphology (RMS = 3.77 nm) with some small “holes” and aggregates, which

### Table 2 Summary of TFT device performances

<table>
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<tr>
<th>Sample/annealing temp.</th>
<th>µh [µh_max]b (cm² V⁻¹ s⁻¹)</th>
<th>Vth (V)</th>
<th>Ion/Ioff</th>
</tr>
</thead>
<tbody>
<tr>
<td>PCZT (N/A)</td>
<td>1.4 × 10⁻⁴ [1.7 × 10⁻⁴]</td>
<td>13</td>
<td>10⁵–10³</td>
</tr>
<tr>
<td>100 °C</td>
<td>1.1 × 10⁻⁴ [1.3 × 10⁻⁴]</td>
<td>–6</td>
<td>10⁵–10³</td>
</tr>
<tr>
<td>150 °C</td>
<td>3.5 × 10⁻⁵ [4.2 × 10⁻⁵]</td>
<td>–10</td>
<td>10⁵–10³</td>
</tr>
<tr>
<td>250 °C</td>
<td>1.4 × 10⁻⁴ [1.5 × 10⁻⁴]</td>
<td>–8</td>
<td>10⁵–10³</td>
</tr>
<tr>
<td>300 °C</td>
<td>3.0 × 10⁻⁴ [4.2 × 10⁻⁴]</td>
<td>–8</td>
<td>10⁵–10³</td>
</tr>
<tr>
<td>PCZTT (N/A)</td>
<td>5.1 × 10⁻⁵ [5.3 × 10⁻⁵]</td>
<td>–7</td>
<td>10⁻⁵–10⁰</td>
</tr>
<tr>
<td>150 °C</td>
<td>1.8 × 10⁻⁴ [1.9 × 10⁻⁴]</td>
<td>–15</td>
<td>10⁻⁵–10⁰</td>
</tr>
<tr>
<td>250 °C</td>
<td>4.8 × 10⁻⁴ [5.1 × 10⁻⁴]</td>
<td>–30</td>
<td>10⁻⁵–10⁰</td>
</tr>
<tr>
<td>300 °C</td>
<td>1.4 × 10⁻³ [1.8 × 10⁻³]</td>
<td>–6</td>
<td>10⁻⁵–10⁶</td>
</tr>
</tbody>
</table>

b Mobility values are measured in air. The mobilities under optimized annealing conditions are indicated in boldface. The average values are calculated over 3 devices and the highest values are listed in parentheses.

### Characterization of molecular organization and film morphologies

Although the TFT performances of the CZ-based polymers were somewhat correlated with their HOMO levels (PCZTT has a shallower HOMO level and higher µh than PCZT), it is necessary to investigate the organization of the thin films by GIWAXS measurements. The 2D-GIWAXS patterns provided information about the molecular packing structures (Fig. 5). The corresponding 1D profiles are shown in Fig. S7.† The pristine thin film of PCZT displayed very weak reflections (Fig. S7a† and S7a†). Only after thermal annealing at 300 °C, one could observe a primary reflection peak (100) located at 2θ = 4.0° along the qy direction (in-plane direction). In addition, a reflection peak (010) located at 2θ = 25.7° along the qx direction (out-of-plane) was found in its 1D profile (Fig. S7b†). This result indicated that PCZT tends to adopt a face-on backbone packing orientation with the chain-to-chain and π-π stacking distances of 22.0 Å and 3.5 Å, respectively (Fig. S8a†,13,25 On the other hand, the thin film of PCZTT exhibited a primary diffraction peak [100] located at 2θ = 4.9° along the qz direction (out-of-plane) with a chain-to-chain packing distance of 18.0 Å (Fig. 5c and S7c†). This result suggested that PCZTT tends to adopt an edge-on backbone packing orientation (Fig. S8b†). Furthermore, the thermal annealing treatment at 300 °C explicitly intensified the primary diffraction peak (Fig. 5d and S7d†). Therefore, the higher mobility of PCZTT was reasonably attributed to its edge-on backbone packing orientations toward the Si substrate, which was favorable for the in-plane charge transport in the TFT devices (Fig. S8b†).10,13,14,22,25 Overall, the GIWAXS results were well correlated with the TFT performances, and it was revealed that the polymers with a planar backbone show a better charge carrier transport property than those with a twist backbone.10,13,13,22,25

![Fig. 5](image-url) GiWAXS patterns of the thin films at different annealing temperatures. PCZT: (a) RT and (b) 300 °C; PCZTT: (c) RT and (d) 300 °C.

![Fig. 6](image-url) Tapping-mode AFM topography images of the thin films of (a to c) PCZT and (d to f) PCZTT, prepared by thermal annealing at different temperatures. Annealing temperatures (for 10 min) are depicted in each image. AFM size: 2 × 2 μm².
indicating an inferior interconnectivity and a low quality thin film [Fig. 6d]. In contrast, much smoother surface morphologies (RMS = 1.52–1.64 nm) with the interconnected grains were obtained after thermal annealing at 150–300 °C (Fig. 6e and f). As a consequence, largely interconnected grains would facilitate the charge carrier transport of the PCZT-based TFTs. Overall, the results were consistent with the observed changes in $\mu_h$ upon thermal annealing.

**Conclusion**

In summary, a new linear-type dibromocarbazole-dioxazine derivative was synthesized and fully characterized. By using this building block as a bifunctional monomer, two CZ-based semiconducting polymers, namely PCZT and PCZTT, were, for the first time, synthesized by Stille polycondensation. The LUMO levels of both polymers were almost the same due to the planarity of both backbone and side chain engineering are currently underway. The charge-transporting properties, estimated from the TFT performances, suggested that PCZT showed a four times higher hole mobility than PCZT. In addition to the HOMO level, the enhanced hole mobility was due to the edge-on backbone packing orientations toward the Si substrate, originating from the stronger intermolecular interactions induced by the planar backbone of PCZTT. This study highlights the promising use of the CZ-based semiconductors in the field of organic electronics. Further improvements of the CZ-based organic semiconductors by backbone and side chain engineering are currently underway in our laboratory.

**Conflicts of interest**

There are no conflicts to declare.

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**Notes and references**
