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The effect of the donor moiety of DPP based polymers on the performance of organic electrochemical transistors†

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Organic mixed (ionic and electronic) charge conductors are the building blocks of state-of-the-art bioelectronic devices, including the organic electrochemical transistors (OECTs). Despite the interest in OECTs, the library of polymers that show efficient mixed charge transport is still narrow. In this work, we developed two donor–acceptor (D–A) type polymers based on the glycolated thiophene diketopyrrolopyrrole (TDPP) as the acceptor unit. We combined the acceptor with two donor units distinguished with different electron-donating strengths, *i.e.*, a glycolated thienylenevinylene (gTVT) and a glycolated thiophene–benzothiadiazole–thiophene (gTBTT), leading to the polymers **TDPP-gTVT** and **TDPP-gTBTT**, respectively. Using spectral, gravimetric, structural, and electrical characterization techniques combined with density functional theory calculations, we sought to understand the effect of the donor unit on the mixed conduction performance of these D–A type polymers. We found that the stronger electron-donating nature of gTVT vs. gTBTT endowed **TDPP-gTVT** with a lower backbone curvature, leading to films with a tighter lamellar packing, and thereof, a higher degree of crystallinity compared to **TDPP-gTBTT**. The combination of a high product of electronic mobility and volumetric capacitance ($\mu C^* = 205 \text{ F cm}^{-1} \text{ V}^{-1} \text{ s}^{-1}$) and a low threshold voltage (-0.36 V) rendered **TDPP-gTVT** based OECTs superior to **TDPP-gTBTT** devices. Our work introduces the selection of donor units with higher electron-donating power as a means to tune the film’s microstructure, providing an effective approach to optimize mixed conduction properties of D–A type OECT materials.

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Introduction

Organic mixed ionic-electronic conductors (OMIECs) transport ionic charges coupled with electronic ones. Mixed conductivity rendered them ideal for developing devices that have an electrolyte interface, such as biosensors, bioactuators, neuromorphic circuits, energy

storage, or conversion devices.^{1,2} A state-of-the-art bioelectronic device that utilizes an OMIEC at the interface with a biological medium is the organic electrochemical transistor (OECT). Compared with the organic field-effect transistor (OFET), which relies on the interfacial accumulation of charges at the channel/dielectric interface, the charge generation in the channel of the OECT occurs in the whole volume of the OMIEC. The volumetric ion-electron coupling in the OECT channel leads to high device transconductance at low gate voltages, typically below 1 V.³ Leveraging the volumetric ion transport in the channel, the OECT acts as a powerful transducer of biochemical events and electrophysiological signals.^{4–6} Conjugated polymers are typical OECT channel materials due to their relatively intact morphology against ion motion and the possibility of tuning their transport properties by chemical synthesis. Operation in aqueous media however requires the conjugated polymers to have oxidation/reduction potentials within a certain electrochemical window to reduce the possibility of side reactions in water. Despite their high transconductance, OECTs typically suffer from slow switching speeds, on the contrary to OFETs, limiting their use in detecting fast-occurring events. However, with proper

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optimization of mixed conductivity, conjugated polymers are promising for high performance and stable OECTs, sought after for real-world applications.⁷

The steady-state performance of OECTs is evaluated in terms of their transconductance (g_m) at saturation conditions, expressed as follows:^{3,8}

$$g_m = \mu C^* \frac{Wd}{L} (V_{TH} - V_G) \quad (1)$$

where W , L , and d are the channel width, length, and thickness, respectively; μ is the charge-carrier mobility and C^* is the capacitance per unit volume of the channel; V_{TH} is the threshold voltage, and V_G is the gate voltage. Eqn (1) shows that the g_m depends on both material and device geometry-related terms. The product of μ and C^* , $[\mu C^*]$, is commonly used as a figure-of-merit to benchmark the mixed conduction properties of OMIECs and predict their performance in OECTs.⁸ V_{TH} is also an important parameter as a low V_{TH} stems from an early onset of charge generation and translates into low power consumption and noise during device operation.

So far, most of the OECTs have relied on the conducting polymer poly(3,4-ethylenedioxythiophene) doped with poly(styrene sulfonate) (PEDOT:PSS) due to its high $[\mu C^*]$, stability, and commercial availability.^{8–10} These transistors operate in the depletion mode as PEDOT:PSS is electrically conducting in its pristine state. The application of V_G turns the OECT OFF. The enhancement mode OECTs, on the other hand, operate in the reverse regime as they use materials that are insulating in their pristine state. In the enhancement mode, the device is turned ON upon application of a V_G , which drives dopant ions inside the film to compensate with electronic charges injected from the contacts. The high ON/OFF current ratios achieved with the enhancement-mode devices reduce the background noise, and the zero-level current in the OFF state demands low power.^{10,11} The abrupt increase in channel current with very low gate voltages is ideal for biosensor applications. Consequently, research efforts in the past years have focused on developing conjugated polymers with maximized $[\mu C^*]$ product for high-performance enhancement-mode OECTs. One common design theme involved rigid and planar electron-rich heterocycles in the backbone.^{12–15} Such an architecture has resulted in high electronic charge carrier mobility because of the enhanced molecular interactions and order in the film.^{14,16} To facilitate ion transport, polar side-chains, such as those comprising various lengths of ethylene glycol (EG) units, were grafted on the backbone, which increased the film's hydrophilicity.^{17,18} As it is the case for OFETs, the side-chain engineering and conjugated backbone chemistry are critical for determining the OECT performance as they govern the solubility, crystallinity, and charge transport properties of the films.^{16,19–22} However, unlike the OFETs, where the relationships between the chemical structure, morphology of the film and device performance are mostly well-established^{23,24} such concepts are yet to be extended to OECTs.^{25,26}

An electron-deficient building block with an easily modifiable skeleton, widely used in OFET and organic photovoltaic devices, is the diketopyrrolopyrrole (DPP).^{9,27–31} DPPs, especially thiophene flanked DPPs (TDPPs), owe their popularity to

the high OFET mobilities (up to $10 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) of their films stemming from their densely π - π stacked units and planar backbones.^{32,33} The backbone planarity of the DPP-based films could be further tuned by attaching various flanked, electron-donating aromatic units.²⁸ Despite the promise of DPPs in OECT applications, only a few examples exist for the DPPs developed as OMIECs. Du *et al.* reported a DPP polymer functionalized with lysine side chains.³⁴ These side chains improved the redox properties of the polymer films and enhanced neural cell adhesion. Reynolds *et al.* reported OECTs comprised of a DPP based polymer with cleavable substituents.³⁵ The polymer was printed from green solvents, and the resulting devices yielded a μC^* of $22 \text{ F cm}^{-1} \text{ V}^{-1} \text{ s}^{-1}$. Giovannitti *et al.* designed pyridine-flanked DPP polymers with a large ionization potential.³⁶ Although these polymers avoided side reactions with oxygen during the device operation, they showed a low electronic charge carrier mobility. McCulloch *et al.* have recently reported three donor-acceptor (D-A) type glycolated TDPPs and highlighted the relationship between the polaron delocalization and the OECT mobility. One of these DPP polymers had a high μC^* of about $300 \text{ F cm}^{-1} \text{ V}^{-1} \text{ s}^{-1}$ but a high threshold voltage of 0.52 V .^{37–41}

Thienylenevinylene (TVT) and thiophene–benzothiadiazole–thiophene (TBTT) are two popular electron-donating units used to form D-A type polymers with high charge-carrier mobilities in OFETs.^{42–44} Incorporating the highly π -extended TVT and TBTT units in the main chain with DPP promotes intermolecular π - π stacking.^{42,43} These two donors have different electron-donating strengths, which are known to influence their interactions with the acceptor, and consequently, the backbone conformation, intramolecular interactions, and the packing behavior of the chains.^{42–45} Motivated by these distinct properties that the donor unit can introduce to the D-A type polymer films,⁴⁴ we developed two new electron-donating moieties, *i.e.*, the glycolated thiophene-vinyl-thiophene (gTVT) and thiophene–benzothiadiazole–thiophene (gTBTT), to construct high-performance OMIECs. We combined these units with a glycolated TDPP and generated two novel D-A type semiconducting polymers. We grafted a suitable length of EG chains on each monomer unit, which improved the polymers' solubility in common organic solvents and endowed the film with redox activity in aqueous electrolytes. By comparing the properties of the two polymers using a combination of solid-state and electrochemical techniques, we systematically elucidated the impact of backbone engineering on the polymer chain conformation, film microstructure, water uptake, surface morphology, and OECT performance. We found that we can tune the backbone curvature *via* the choice of the donor unit, which enabled us to develop an OMIEC with a low threshold voltage (-0.36 V) and high OECT performance (μC^* of $205 \text{ F cm}^{-1} \text{ V}^{-1} \text{ s}^{-1}$) competing with that of the PEDOT:PSS-based devices.

Results and discussion

Synthesis

Two novel monomers of bis(trimethylstannylyl)-TVT and bis(trimethylstannylyl)-TBTT were synthesized, and their chemical



structures and intermediate products (**1–6**) are shown in Scheme 1. The bis(trimethylstannyl)-TVT **4** begins with nucleophilic substitution of 3-bromothiophene to obtain the glycolated-thiophene **1**, followed by the bromination with NBS at low temperature, giving glycolated 2-bromo thiophene **2**.⁴⁶ A Stille coupling between **2** and bis(tributylstannyl)ethene afforded the glycolated-TVT unit (**3**) and then stannization reaction produced the key monomer **4**. Bis(trimethylstannyl)-TBTT **6** was synthesized by methods similar to **4**,^{46,47} except that we used the reagent 4,7-bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2yl)2,1,3-benzothiadiazole. We chose to have two EG units on the side chains to enhance the polymer's solubility in common organic solvents and aid in the uptake of hydrated ions of the electrolyte without causing dissolution or delamination of the films in water.

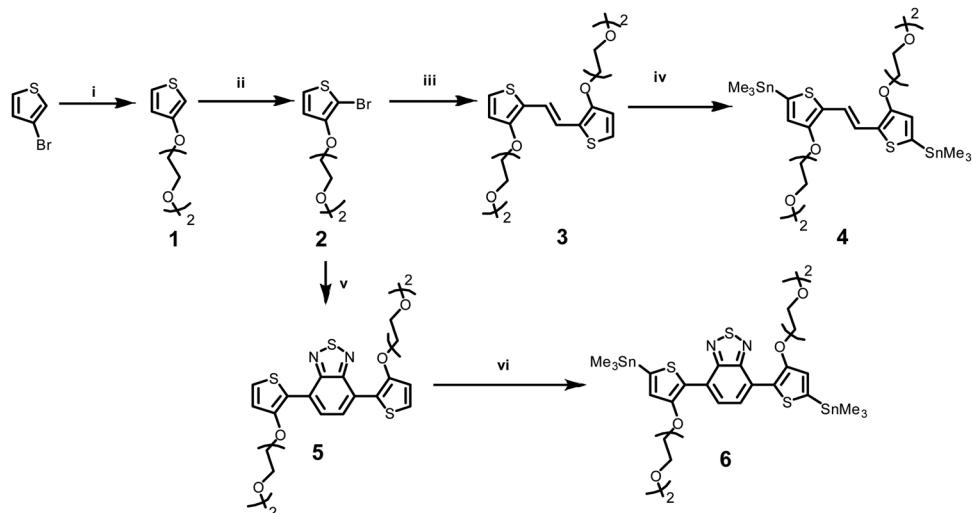
Besides functionalizing the electron-donating monomers with EG units, we incorporated linear EG chains ($n = 4$) to the DPP with the anticipation to enhance the films' capacitance (Scheme 2). Tosylation and subsequent glycolation with DPP afforded the non-halogenated thiophene DPP, which was finally brominated to give our acceptor unit. Stille polymerization of the glycolated thiophene dibromoDPP and 2,5-bis(trimethylstannyl)-TVT or 2,5-bis(trimethylstannyl)-TBTT was then used to generate the two polymers, namely, **TDPP-gTVT** and **TDPP-gTBTT**, in high yields (Scheme 2). The linear oligo EG side-chains of TVT and TBTT had the same lengths. Like the EG functionalized polymers reported before,⁴⁷ the polymers' high polarity did not allow the molecular weight analysis by gel permeation chromatography (GPC) in solvents such as chlorobenzene, chloroform, or dimethylformamide. We, therefore, used the matrix-assisted laser desorption/ionization time-of-flight spectrometry (MALDI-TOF) and estimated comparable molecular weights ranging from 4.0 kDa to 8.0 kDa (Fig. S1, ESI†). The resulting polymers showed excellent

solubility in common solvents, such as dichloromethane and chloroform, ($\approx 10 \text{ mg mL}^{-1}$). The materials also exhibited good thermal stability with the degradation temperature above 340 °C (Fig. S2, ESI†).

Optoelectrical properties

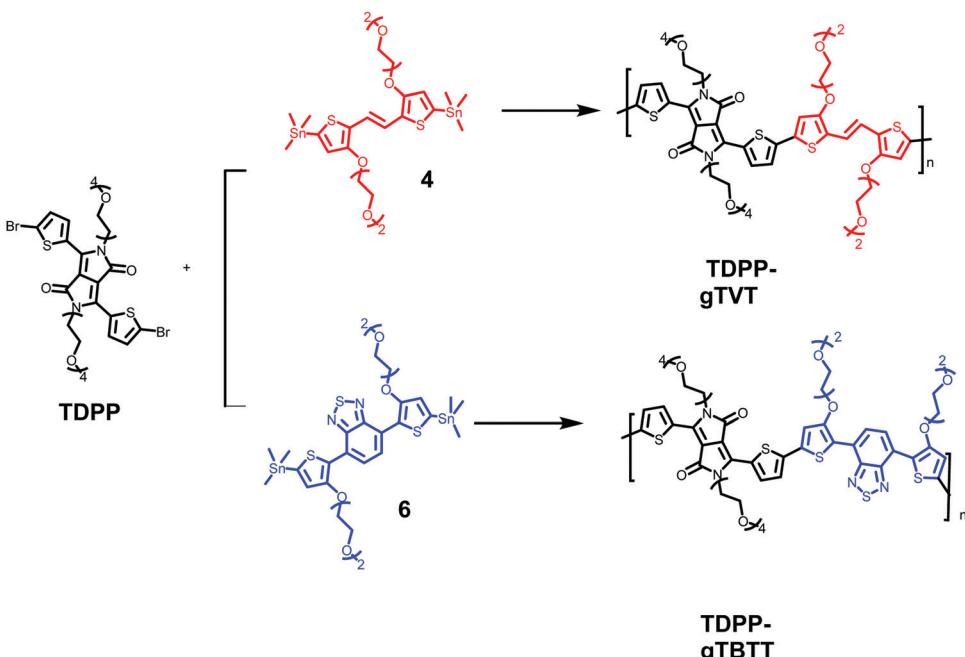
To study the effect of backbone chemistry on the polymers' aggregation properties, we recorded the UV-Vis-NIR absorption spectra of conjugated polymers in solution and as thin films. All the absorption spectra show a similar profile with dual-band absorption (see Fig. 1a for the thin film spectra and Fig. S3 (ESI†) for the solution spectra). We observe a high energy band spanning from 350 nm to 500 nm in all spectra, attributed to the $\pi-\pi^*$ transition. The low-energy bands originate from an intramolecular charge transfer (ICT) between the DPP and the donors. In solution, we record an absorption maximum (λ_{max}) at 853 nm for **TDPP-gTVT**, and 785 nm for **TDPP-gTBTT** (Table 1). When going from the solution to the thin film, the spectra of both films display a red-shift, *i.e.*, the λ_{max} of **TDPP-gTVT** and **TDPP-gTBTT** shifts to 878 nm and 813 nm, respectively, indicating that some degree of aggregation or $\pi-\pi$ stacking occurs in the solid-state.⁴⁸ Notably, while the change in λ_{max} when going from the solution-to-thin film is larger for **TDPP-gTBTT**, the absorption maximum is at higher wavelengths for **TDPP-gTVT**, both in solid and solution state. The red-shifted absorption of **TDPP-gTVT** stems from the stronger donating strength due to the more electron-rich nature of gTVT compared to gTBTT. This trend is consistent with their all-alkylated polymer analogues.^{42,45}

To elucidate the effect of the donor unit on the electron density distribution and energetics, we carried out density functional theory (DFT) calculations on methyl-substituted DPP and methoxy-substituted gTVT and gTBTT of **TDPP-gTVT**



Scheme 1 The synthetic route of bis(trimethylstannyl) glycolated TVT, bis(trimethylstannyl) glycolated TBTT monomers. Reaction conditions: (i) *t*-BuOK, 2-(2-methoxyethoxy)ethanol, RT, then 100 °C; (ii) NBS, THF/Hexane, 0 °C; (iii) 2-bis(tributylstannyl)ethene, $\text{Pd}_2(\text{dba})_3$, $\text{P}(\text{o-tol})_3$, 100 °C; (iv) *n*-BuLi, THF, −78 °C, SnClMe_3 , then RT; (v) 4,7-Bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2yl)2,1,3-benzothiadiazole, $\text{Pd}_2(\text{dba})_3$, $\text{P}(\text{o-tol})_3$, tricaprylylmethylammonium chloride, aq. K_3PO_4 , toluene; (vi) *n*-BuLi, THF, −78 °C, SnClMe_3 , then RT.





Scheme 2 The synthetic route of the glycolated DPP polymers: TDPP-gTVT and TDPP-gTBTT.

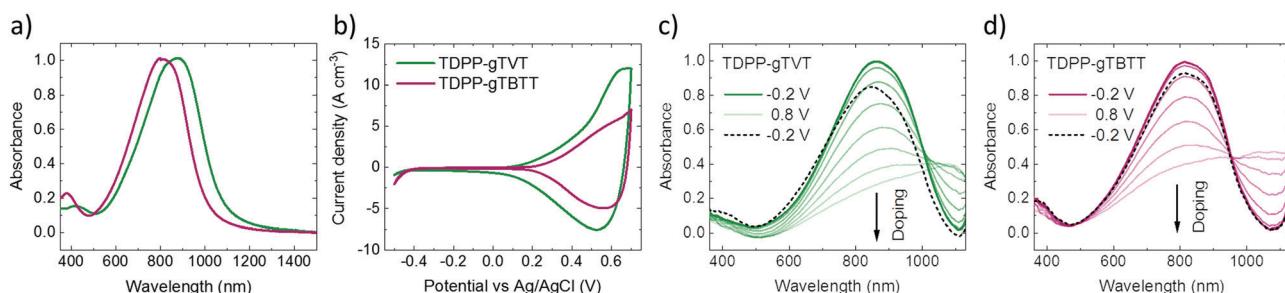


Fig. 1 (a) Normalized UV-Vis-NIR absorption spectra and (b) cyclic voltammograms of the polymer films spin-cast from chloroform on ITO substrates. The CV curves were recorded with a scanning speed of 80 mV s^{-1} aqueous NaCl solution (0.1 M). The 10th cycle is shown. (c and d) UV-Vis absorption spectra of the films during the forward doping sweep (-0.2 V to 0.8 V) direction and as they were de-doped (black dashed spectrum, at -0.2 V). All measurements were performed in aqueous NaCl solution (0.1 M) and voltages were applied vs. Ag/AgCl. The film thickness was in the range of 80–125 nm.

Table 1 Optoelectrical and thermal properties of the polymers

Polymer	λ_{soln}^a [nm]	λ_{film}^b [nm]	IP ^c [eV]	EA ^d [eV]	E_{ox}^e [V]	E_{opt}^f [eV]
TDPP-gTVT	853	878	4.44	3.30	0.21	1.14
TDPP-gTBTT	785	812	4.60	3.37	0.29	1.23

^a In chloroform. ^b As thin films. ^c Ionization potentials (IPs) were estimated from the onset of the first oxidation peak (E_{ox} vs. Fc/Fc^+) in $n\text{-Bu}_4\text{NPF}_6$ using the equation $\text{IP}(\text{eV}) = 4.8 - E_{\text{ox}}$. ^d Electron affinities (EAs) were estimated according to $\text{EA}(\text{eV}) = \text{IP} - E_{\text{opt}}$. ^e Estimated from the onset of the first oxidation peak in 0.1 M aqueous NaCl. ^f Determined from the absorption onset in thin-film UV-Vis-NIR spectra.

and TDPP-gTBT by Gaussian 09 with the level of B3LYP/6-31G. The geometry optimized structures reveal that each trimer approaches planarity with very small inter-ring torsion

(Fig. S4, ESI[†]). However, the backbone curvature based on the through-space distance is distinctive across the two polymers (Fig. S5, ESI[†]). Our calculations show that the degree of curvature of the conjugated backbone differs depending on the donor unit, with the dihedral angle between the comonomers in the following order: TDPP-gTVT (157°) > TDPP-gTBTT (134°). We expect the TDPP-gTVT with lower backbone curvature to exhibit stronger interchain interactions, leading to a higher degree of crystallinity.³³

We next extracted the energy levels of our polymer films using cyclic voltammetry (CV) measurements performed in 0.1 M tetrabutylammonium hexafluorophosphate ($n\text{-Bu}_4\text{NPF}_6$) in acetonitrile (Fig. S6, ESI[†]). The highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) energy levels are -4.44 eV and -3.30 eV for TDPP-gTVT, and -4.60 eV and -3.37 eV for TDPP-gTBTT (Table 1).

The simulated HOMO/LUMO levels are -4.28 eV and -2.88 eV for **TDPP-gTBT**, and -4.61 eV and -3.16 eV for **TDPP-gTBTT** (Fig. S5, ESI[†]). The trend of the calculated HOMO and LUMO is consistent with experimental results, *i.e.*, both simulated levels of **TDPP-gTBT** are higher than those of **TDPP-gTBTT**. As expected from the electron-withdrawing property of BT, **TDPP-gTBTT** shows lower HOMO and LUMO values than the **TDPP-TBT** that bears the vinylene motif. These results agree well with the DFT calculations summarized in Fig. S5 (ESI[†]). Our calculations suggest that HOMO of both polymers is delocalized over the backbone while LUMO localizations differ. The LUMO of **TDPP-gTBT** is mainly located on the TDPP and the neighboring thiophene of the TBT. As for **TDPP-gTBTT**, LUMOs are located on the TDPP, the adjacent thiophene, as well as the BT moiety of the TBTT. Therefore, while the HOMO levels of the two polymers are affected by both donor and acceptor units, the LUMO level of **TDPP-gTBTT** is affected more by the acceptor unit, which can be ascribed to the electron-accepting nature of BT. Finally, the high HOMO levels of these polymers suggest that they are promising to build p-type enhancement-mode OECTs.

Electrochemical analysis and OECT characterization

We recorded the CV curves of the films in 0.1 M NaCl solution to characterize the extent of electrochemical switching between the neutral and oxidized states (Fig. 1b). Both polymers show an increase in the current in the p-type region with a low onset of oxidation below 0.3 V and redox peaks appearing at *ca.* 0.5 V *versus* the reference electrode (Ag/AgCl). The polymers show electrochemical stability, deduced from their relatively constant current profiles over ten CV cycles (Fig. S7, ESI[†]). The gTBT analog exhibits a lower onset (0.21 V *vs.* 0.29 V) and higher oxidation currents than gTBTT. Fig. 1c and d show the UV-VIS absorption spectrum changes as the polymer films are electrochemically doped in 0.1 M NaCl. We see that, for both films, when they are biased at positive voltages *versus* Ag/AgCl, the broad absorption band extending from 600 nm to 1000 nm is bleached, while a new band appears above 1000 nm. We attribute the latter to the formation of polarons as anions are injected into the films. Neither of the spectra, however, seem to fully recover upon de-doping at a bias of opposite polarity

(*i.e.*, -0.2 V), while the electrochromic property of **TDPP-gTBTT** appears to be slightly more reversible than **TDPP-TBT**.

As the films showed promising HOMO levels and electrochemical doping performance in the aqueous electrolyte, we next fabricated micron-scale OECTs to assess the effect of the donor units on device performance. The films were patterned on OECT channels that had identical geometry ($W = 100$ μm , $L = 10$ μm). The OECT output, transfer and transconductance curves are shown in Fig. 2. While both films switched ON upon application of a negative V_G , **TDPP-gTBT** OECTs showed much better device characteristics. For instance, **TDPP-TBT** OECTs generated about 4-fold higher ON-currents than the **TDPP-gTBTT** devices and exhibited a geometry normalized transconductance, $g_m/(Wd/L)$, of 47.8 ± 2.8 S cm^{-1} , at least one order of magnitude higher than the **TDPP-gTBTT** devices (3.65 ± 0.13 S cm^{-1}). **TDPP-gTBT** has also a lower threshold voltage ($V_{TH} = -0.36$ V *vs.* $V_{TH} = -0.42$ V). The ON/OFF ratios measured are in the order of 10^4 – 10^5 , which is at the same range with the recently reported D-A type OECT polymers.^{34–37} Using eqn (1) above and the steady-state characteristics recorded, we found that **TDPP-gTBT** has a higher $[\mu\text{C}^*]$ of 205.2 ± 12 $\text{F cm}^{-1} \text{V}^{-1} \text{s}^{-1}$ than **TDPP-gTBTT** ($[\mu\text{C}^*]$ of 21.5 ± 0.7 $\text{F cm}^{-1} \text{V}^{-1} \text{s}^{-1}$). The steady-state performance of **TDPP-gTBT** OECTs is comparable to the performance of other recently published high-performance D-A polymers or all donor polymers.^{34–37} Importantly, the top OECT performer in this work showed a lower threshold voltage ($V_{TH} = -0.36$ V) than those of the previously reported D-A polymers, such as *p*(gDPP-T2) ($V_{TH} = -0.52$ V) and **PDPP-DT** ($V_{TH} = -0.59$ V) (Table S1, ESI[†]).^{37,49} When we recorded the transient behaviour of these devices, we found a response time of around 7.3 ms and 8.7 ms for **TDPP-gTBT** and **TDPP-gTBTT**, respectively (Fig. S8a and b, ESI[†]), indicating that the **TBT** variant has a faster response than **TBTT**. The devices could however not withstand to the application of square shaped voltage pulses at the gate electrode, *i.e.*, the I_D decreased over time (Fig. S8c and d, ESI[†]). **TDPP-gTBTT** devices were more stable than **TDPP-gTBT**, which can also be linked to higher electrochromic reversibility of the **TBTT** variant (Fig. 1c and d). Note, however, that CV curves of both polymers showed only a negligible change upon

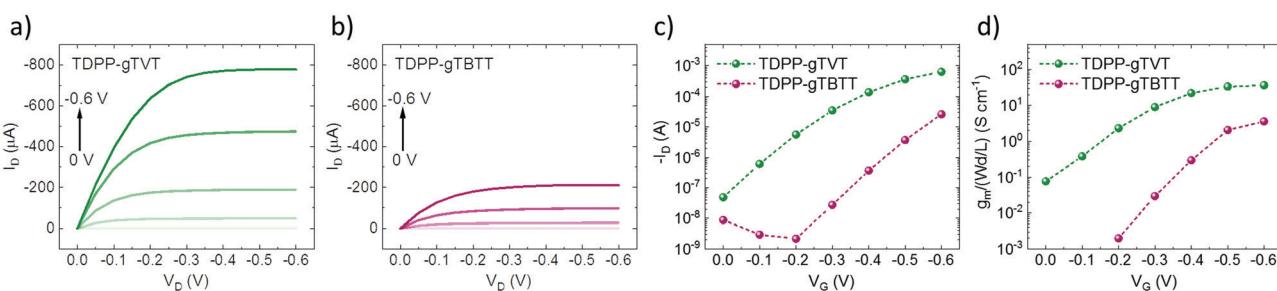


Fig. 2 OECT output (a and b), transfer (c) and transconductance curves (d) curves. V_G changed from 0 V to -0.6 V with a step of -0.1 V. Transconductance (g_m) values were calculated at a drain voltage of $V_D = -0.6$ V and normalized with channel dimensions. The gate electrode was Ag/AgCl and NaCl (0.1 M, aqueous) was used as the electrolyte. The channels had a width of 100 μm and a length of 10 μm . The channel thicknesses were 73 nm and 45 nm, for **TDPP-gTBT** and **TDPP-gTBTT**, respectively.



cycling (Fig. S6 and S7, ESI[†]). This discrepancy can be explained by the fact that in OECTs, the electronic conductivity in the channel is the first to suffer from the side effects of swelling, as interconnectivity between electronically conductive regions may become disrupted due to a morphology change upon ion/water uptake. The CV profile may be less affected by such morphological changes as the current response is not limited to horizontal charge transport and governed fully by the interconnectivity of chains. It is also the case that films with higher degree of aggregation tend to be more susceptible to electronic disruption due to electrolyte uptake,⁵⁰ which can explain why the **TDPP-gTVT** OECT appears to break down faster than that of **TDPP-gTBTT**. To prevent this degradation and obtain OECTs robust against pulsing, one may consider increasing the conjugated polymer length, while retaining the film's characteristic electrochemical properties.

To understand which polymer property governs the OECT behavior and how it is affected by the donor unit in the structure, we investigated our materials' ion uptake and transport abilities. We, first, recorded the electrochemical impedance spectra of these films (Fig. S9 and S10, ESI[†]). As the films were electrochemically doped at 0.6 V vs. Ag/AgCl, the impedance magnitude decreased, and the phase gained a capacitive character for two polymers. The data in the doped state could be fit with $R_s(R_p||C)$ equivalent circuit, allowing us to extract the capacitance values of each film (Fig. S11, ESI[†]). The C^* values at low frequencies exhibit a perfect plateau (Fig. S11, ESI[†]), and C^* at $f = 0.1$ Hz agree with those calculated from the equivalent circuit fit. **TDPP-gTVT** has a C^* of *ca.* 173.5 F cm⁻³, higher than that of **TDPP-gTBTT** (122.5 F cm⁻³). Calculating the C^* allows to estimate the electronic mobility of each polymer in the OECT channel (μ_{OECT}). The μ_{OECT} of **TDPP-gTVT** is about an order of magnitude higher than **TDPP-gTBTT** (*ca.* 1.1 cm⁻¹ V⁻¹ s⁻¹ vs. 0.18 cm⁻¹ V⁻¹ s⁻¹). Since the C^* values are relatively comparable, these results show that the high performance of **TDPP-gTVT** in OECTs stems mainly from the film's high charge carrier mobility. We summarize the OECT characteristics and the materials figures of merit in Table 2.

It is interesting to note that the CV curves, the polymer energetics or the spectroelectrochemistry data do not predict such a large difference in the OECT performance of these two polymers. To elucidate the origin of the high performance of TVT unit *versus* TBTT, we performed a combination of morphological and structural studies. First, we used electrochemical quartz crystal microbalance with dissipation monitoring (EQCM-D) technique to evaluate whether these differences are

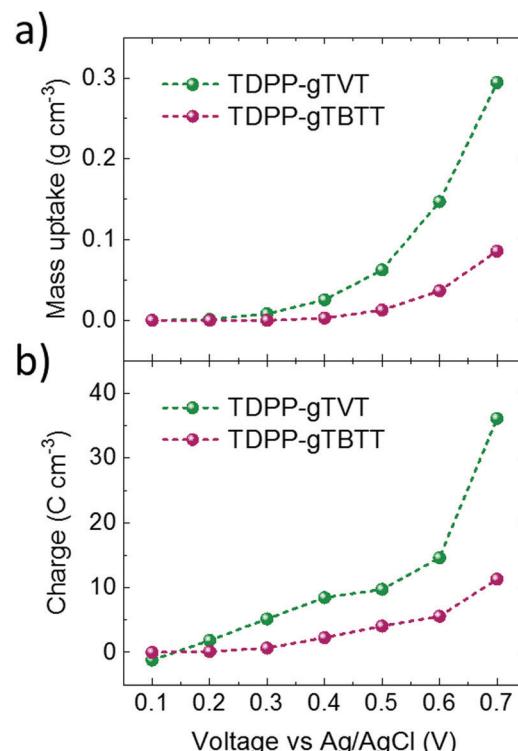


Fig. 3 Electrochemical quartz crystal microbalance with dissipation monitoring (EQCM-D) results of **TDPP-gTVT** and **TDPP-gTBTT**. The differences in (a) the electrolyte uptake and (b) charge generation per unit volume during the application of doping potentials from 0.1 V to 0.7 V vs. Ag/AgCl with a step of 0.1 V.

related to the electrolyte uptake behavior of the polymer films. EQCM-D measured the mass accumulating inside the polymer films as they were subject to doping potentials. We observe an initial mass increase for both films upon exposure to the electrolyte due to passive film swelling. **TDPP-gTVT**, however, exhibits passive swelling of nearly double that of **TDPP-gTBTT** (22.5% vs. 13.2%, respectively (Fig. S12, ESI[†])). As we start applying doping potentials, both films start gaining mass. Fig. 3a shows that **TDPP-gTVT** exhibits an earlier onset of electrolyte uptake into the film (0.3 V) compared to **TDPP-gTBTT** (0.5 V) and takes up more than 3 times the amount of electrolyte at a doping potential of 0.7 V. The enhanced electrolyte uptake of the **TDPP-gTVT** film is accompanied with an earlier and greater charge generation (Fig. 3b). The charging behavior in **TDPP-gTVT** is thus in line with its high C^* . As the film swells more readily, ion penetration upon

Table 2 Steady-state characteristics of OECTs comprising **TDPP-gTVT** and **TDPP-gTBTT**

Polymer	$I_{\text{D ON:OFF}}$	V_{TH} (V)	μ_{OECT}^a (cm ² V ⁻¹ s ⁻¹)	$C^* b$ (F cm ⁻³)	$\mu C^* c$ (F cm ⁻¹ V ⁻¹ s ⁻¹)	g_m^d (Wd L ⁻¹) ^d (S cm ⁻¹)	I_{G} leakage current ^e (nA)
TDPP-gTVT	10^5	-0.36	1.1 ± 0.07	173.5 ± 5	205.2 ± 12	47.8 ± 2.80	-89.7
TDPP-gTBTT	10^5	-0.42	0.18 ± 0.006	122.5 ± 15	21.5 ± 0.7	3.65 ± 0.13	-28.8

^a OECT hole mobility of the films calculated using eqn (1) by using the C^* values. ^b Calculated from the electrochemical impedance spectra using $R_s(R_p||C)$ fits.

^c Estimated using the eqn (1), at maximum g_m . ^d The highest g_m values normalized with the channel thickness. Error bars for the capacitance measurements arise from thickness variations. For OECTs measurements, at least 3 channels were measured. ^e Gate current recorded at $V_G = V_D = -0.6$ V.



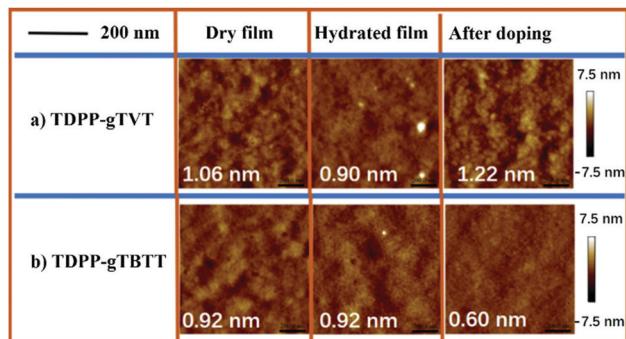


Fig. 4 Atomic force microscopy (AFM) height images of (a) TDPP-gTVT and (b) TDPP-gTBTT, along with the RMS values in dry (as-cast), electrolyte swollen and after being subject to two oxidative CV cycles.

biasing is easier, promoting a greater extent of ionic-electronic charge coupling.

To understand what kind of film morphology leads to the improved electrolyte uptake, C^* and μ_{OECT} , we next performed tapping-mode atomic force microscopy (AFM). AFM scanned the overall surface morphology of the films in their as-cast, hydrated in 0.1 NaCl solution, and after two oxidative CV cycles (Fig. 4). AFM images show that both films have nanofibrillar intercalated structures in their pristine state. The root-mean-square roughness (RMS) of the films is also similar: TDPP-gTVT shows an RMS of 1.06 nm and TDPP-gTBTT has an RMS of 0.92 nm. However, the surface morphology of TDPP-gTVT

changes significantly in the hydrated state with a reduction in the RMS (0.90 nm), in contrast to the TDPP-gTBTT film, which shows negligible changes when going from a dry-to-swollen state. The smoothed surface is indicative of a film swollen by ions and water. After doping–dedoping cycles were completed, TDPP-gTVT exhibited fibrillar features characterized by a much rougher height profile (RMS = 1.22 nm). TDPP-gTBTT, on the other hand, became smoother, evidenced by a low RMS of 0.60 nm. We argue that the surface becomes rougher because of ion and water penetration and ejection cycles upon doping and de-doping.⁵¹ Formation of distinct regions with different textures is a sign of ion movement in and out of the film, in agreement with the EQCM-D findings.

Ex situ cycled and swollen state AFM images evidence the changes TDPP-gTVT undergoes due to effective electrochemical doping. However, since the film morphologies of the two films are very similar in their pristine state, AFM does not provide any insight on why one polymer outperforms the other in OECTs. To investigate the packing behavior of chains, we performed two-dimensional grazing incident wide-angle X-ray scattering (GIWAXS) studies (Fig. 5, see the packing parameters summarized in Table S2, ESI[†]). Scattering profiles show that TDPP-gTVT and TDPP-gTBTT exhibit a lamellar diffraction peak (100) along the in-plane direction at $q_{xy} = 0.332$ and 0.329 \AA^{-1} , which corresponds to a lamellar d -spacing of 18.92 and 19.09 \AA , respectively. The shorter lamellar stacking distance of TDPP-gTVT suggests a tighter packing in the lamellar

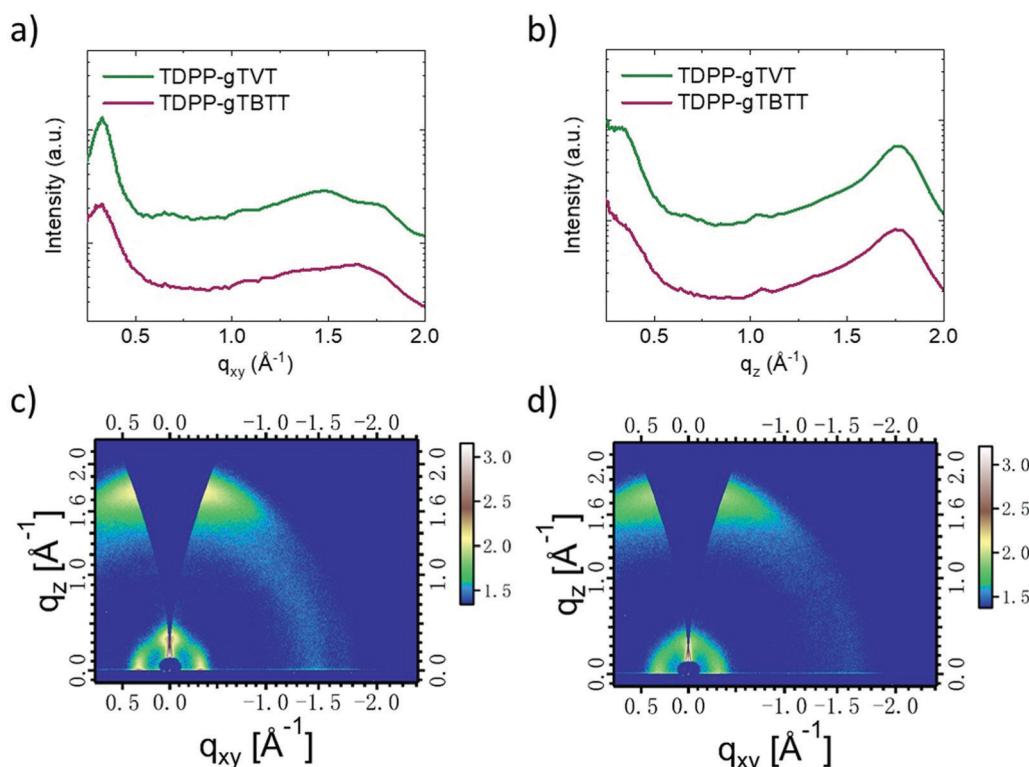


Fig. 5 Line-cut profiles of TDPP-gTVT and TDPP-gTBTT films obtained by integration along the (a) in-plane (q_{xy}) and (b) out-of-plane (q_z) direction. 2D grazing-incidence wide-angle X-ray scattering (GIWAXS) images of (c) TDPP-gTVT and (d) TDPP-gTBTT films; q_z and q_{xy} are the perpendicular and parallel wave vector transfers with respect to the substrate surface.



direction. **TDPP-gTBT** and **TDPP-gTBTT** also exhibit a significant out-of-plane (010) π - π scattering peak at $q_z = 1.75$ and 1.73 \AA^{-1} , corresponding to a π spacing of 3.59 and 3.63 \AA , respectively. Together with the lamellar diffraction peak (100) peak in the in-plane direction, this peak suggests a predominantly face-on orientation where π - π stacks are perpendicular to the substrate. Using (010) diffractions, we also calculated the crystal coherence lengths (CCLs) according to the Scherrer equation. We summarized these values in Table S2 (ESI[†]). The CCL_{OOP,010} of **TDPP-gTBT** in the out-of-plane direction is estimated to be 23.5 \AA , larger than **TDPP-gTBTT** (20.2 \AA). The larger CCL of **TDPP-gTBT** suggests higher crystallinity.

Although **TDPP-gTBT** and **TDPP-gTBTT** form films with similar chain orientation with respect to the substrate, **TDPP-gTBT** exhibits a higher order in the lamellar stacking direction, a denser π -stacking, and longer CCL, resulting in a higher charge carrier mobility. Recalling the red-shifted spectrum of **TDPP-gTBT** (Fig. 1a), we suggest that this film has a microstructure with better connected or larger aggregates. Moreover, the lower backbone curvature of **TDPP-gTBT**, revealed by the DFT calculations, indicates stronger intermolecular interactions, which agrees well with the scattering results. Taken together, these results suggest that by carefully choosing the donor unit, we can fine-tune the packing behavior of the D-A polymer chains through the backbone curvature, which affects their charge-transport ability. Surprisingly, the TBT unit also enables larger swelling upon exposure to electrolyte and electrochemical doping, which leads to higher C^* . For this polymer, mobility, promoted by crystallinity, does not come at the expense of ion/water uptake and high capacitance. We postulate that improving both of the parameters that lead to high-performance OECTs was possible by including EG units on both the donor and the acceptor (which promotes ion uptake) while maintaining a rather rigid backbone (which allows for electronic charge transport). Note also that our polymers have relatively short chains, which help form better-stacked aggregates compared with other OMIECs.

Conclusion

In this study, we developed two novel glycolated DPP based copolymers by introducing gTBT and gTBTT units on the backbone. We examined how the donating strength of the donor unit affects the polymer film optical, morphological, electrochemical, and water swelling properties as well as the steady-state performance in microfabricated OECTs. **TDPP-gTBT** film took up more electrolyte when immersed in it, changed its morphology upon electrochemical doping/dedoping cycles and accumulated more mass upon doping compared to **TDPP-gTBTT**. **TDPP-gTBT** exhibited a red-shifted absorption and denser π -stacking. In OECTs, **TDPP-gTBT** exhibited a relatively high C^* of 173.5 F cm^{-3} , μC^* of $205 \text{ F cm}^{-1} \text{ V}^{-1} \text{ s}^{-1}$ and a low V_{TH} of -0.36 V . The superior performance of **TDPP-gTBT** over **TDPP-gTBTT** is ascribed to (1) the enhanced affinity to the electrolyte and larger amount of

charge generated, (2) the low backbone curvature, which leads to a high degree of crystallinity, promoting high hole carrier mobility. Both polymers however show low device stability, revealed from the decrease in the drain current upon application of voltage pulses at the gate electrode. The instability can be addressed by increasing the molecular weight of the polymers produced. Our results suggest that it is possible to balance capacitance and mobility, which often requires contracting structural features (soft-amorphous content and high crystallinity, respectively) by tuning the donating strength of polymers with minimized backbone curvature and sufficient hydrophilicity.

Author contributions

The manuscript was written through the contributions of all authors. Y. Wang, A. Hamidi-Sakr, and Jokubas Surgailis have contributed equally to this work. Wan Yue and Sahika Inal contributed equally to this work.

Conflicts of interest

The authors declare no competing financial interest.

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References

- 1 B. D. Paulsen, K. Tybrandt, E. Stavrinidou and J. Rivnay, *Nat. Mater.*, 2020, **19**, 13–26.
- 2 D. Ohayon and S. Inal, *Adv. Mater.*, 2020, **32**, 2001439.
- 3 J. Rivnay, S. Inal, A. Salleo, R. M. Owens, M. Berggren and G. G. Malliaras, *Nat. Rev. Mater.*, 2018, **3**, 17086.
- 4 S. Wustoni, C. Combe, D. Ohayon, M. H. Akhtar, I. McCulloch and S. Inal, *Adv. Funct. Mater.*, 2019, **29**, 1904403.
- 5 E. Macchia, P. Romele, K. Manoli, M. Ghittorelli, M. Magliulo, Z. M. Kovács-Vajna, F. Torricelli and L. Torsi, *Anal. Bioanal. Chem.*, 2020, **412**, 5005–5014.
- 6 D. Khodagholy, T. Doublet, P. Quilichini, M. Gurfinkel, P. Leleux, A. Ghestem, E. Ismailova, T. Hervé, S. Sanaur, C. Bernard and G. G. Malliaras, *Nat. Commun.*, 2013, **4**, 1575.
- 7 S. Inal, J. Rivnay, A. Suiu, G. G. Malliaras and I. McCulloch, *Acc. Chem. Res.*, 2018, **51**, 1368–1376.
- 8 S. Inal, G. G. Malliaras and J. Rivnay, *Nat. Commun.*, 2017, **8**, 1767.
- 9 Y. Fu, N. Wang, A. Yang, H. W. Law, L. Li and F. Yan, *Adv. Mater.*, 2017, **29**, 1703787.



10 W. Lee, D. Kim, N. Matsuhisa, M. Nagase, M. Sekino, G. G. Malliaras, T. Yokota and T. Someya, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 10554–10559.

11 A. M. Pappa, D. Ohayon, A. Giovannitti, I. P. Maria, A. Savva, I. Uguz, J. Rivnay, I. McCulloch, R. M. Owens and S. Inal, *Sci. Adv.*, 2018, **4**, eaat0911.

12 E. Zeglio and O. Inganäs, *Adv. Mater.*, 2018, **30**, 1800941.

13 C. B. Nielsen, A. Giovannitti, D. T. Sbircea, E. Bandiello, M. R. Niazi, D. A. Hanif, M. Sessolo, A. Amassian, G. G. Malliaras, J. Rivnay and I. McCulloch, *J. Am. Chem. Soc.*, 2016, **138**, 10252–10259.

14 A. Giovannitti, D. T. Sbircea, S. Inal, C. B. Nielsen, E. Bandiello, D. A. Hanif, M. Sessolo, G. G. Malliaras, I. McCulloch and J. Rivnay, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, 12017–12022.

15 E. Zeglio, J. Eriksson, R. Gabrielsson, N. Solin and O. Inganäs, *Adv. Mater.*, 2017, **29**, 1605787.

16 M. Moser, J. F. Ponder Jr, A. Wadsworth, A. Giovannitti and I. McCulloch, *Adv. Funct. Mater.*, 2019, **29**, 1807033.

17 Y. Wang, E. Zeglio, H. Liao, J. Xu, F. Liu, Z. Li, I. P. Maria, D. Mawad, A. Herland, I. McCulloch and W. Yue, *Chem. Mater.*, 2019, **31**, 9797–9806.

18 Y. Wang, Y. Yu, H. Liao, Y. Zhou, I. McCulloch and W. Yue, *Acc. Chem. Res.*, 2020, **53**, 2855–2868.

19 C. B. Nielsen, M. Turbiez and I. McCulloch, *Adv. Mater.*, 2013, **25**, 1859–1880.

20 J. Mei and Z. Bao, *Chem. Mater.*, 2014, **26**, 604–615.

21 M. Moser, T. C. Hidalgo, J. Surgailis, J. Gladisch, S. Ghosh, R. Sheelamanthula, Q. Thiburce, A. Giovannitti, A. Salleo, N. Gasparini, A. Wadsworth, I. Zozoulenko, M. Berggren, E. Stavrinidou, S. Inal and I. McCulloch, *Adv. Mater.*, 2020, 2002748.

22 P. Schmode, A. Savva, R. Kahl, D. Ohayon, F. Meichsner, O. Dolynchuk, T. Thurn-Albrecht, S. Inal and M. Thelakkat, *ACS Appl. Mater. Interfaces*, 2020, **12**, 13029–13039.

23 E. J. W. Crossland, K. Tremel, F. Fischer, K. Rahimi, G. Reiter, U. Steiner and S. Ludwigs, *Adv. Mater.*, 2012, **24**, 839–844.

24 A. Hamidi-Sakr, L. Biniek, S. Fall and M. Brinkmann, *Adv. Funct. Mater.*, 2016, **26**, 408–420.

25 J. W. Onorato and C. B. Luscombe, *Mol. Syst. Des. Eng.*, 2019, **4**, 310–324.

26 J. H. Kim, S. Kim, G. Kim and M. Yoon, *Macromol. Biosci.*, 2020, 2000211.

27 L. R. Savagian, A. M. Österholm, J. F. Ponder Jr., K. J. Barth, J. Rivnay and J. R. Reynolds, *Adv. Mater.*, 2018, 1804647.

28 X. Guo, A. Facchetti and T. J. Marks, *Chem. Rev.*, 2014, **114**, 8943–9021.

29 R. S. Ashraf, I. Meager, M. Nikolka, M. Kirkus, M. Planells, B. C. Schroeder, S. Holliday, M. Hurhangee, C. B. Nielsen, H. Sirringhaus and I. McCulloch, *J. Am. Chem. Soc.*, 2015, **137**, 1314–1321.

30 I. Meager, R. S. Ashraf, S. Mollinger, B. C. Schroeder, H. Bronstein, D. Beatrup, M. S. Vezie, T. Kirchartz, A. Salleo, J. Nelson and I. McCulloch, *J. Am. Chem. Soc.*, 2013, **135**, 11537–11540.

31 Z. Liu, G. Zhang and D. Zhang, *Acc. Chem. Res.*, 2018, **51**, 1422–1432.

32 I. Kang, H.-J. Yun, D. S. Chung, S.-K. Kwon and Y.-H. Kim, *J. Am. Chem. Soc.*, 2013, **135**, 14896–14899.

33 J. Y. Back, H. Yu, I. Song, I. Kang, H. Ahn, T. J. Shin, S.-K. Kwon, J. K. Oh and Y.-H. Kim, *Chem. Mater.*, 2015, **27**, 1732–1739.

34 W. Du, D. Ohayon, C. Combe, L. Mottier, I. P. Maria, R. S. Ashraf, H. Fiumelli, S. Inal and I. McCulloch, *Chem. Mater.*, 2018, **30**, 6164–6172.

35 B. Schmatz, A. W. Lang and J. R. Reynolds, *Adv. Funct. Mater.*, 2019, **29**, 1905266.

36 A. Giovannitti, R. B. Rashid, Q. Thiburce, P. D. Paulsen, C. Cendra, K. Thorley, D. Moia, J. T. Mefford, D. Hanif, D. Weiyuan, M. Moser, A. Salleo, J. Nelson, I. McCulloch and J. Rivnay, *Adv. Mater.*, 2020, **32**, 1908047.

37 M. Moser, A. Savva, K. Thorley, B. D. Paulsen, T. C. Hidalgo, D. Ohayon, H. Chen, A. Giovannitti, A. Marks, N. Gasparini, A. Wadsworth, J. Rivnay, S. Inal and I. McCulloch, *Angew. Chem., Int. Ed.*, 2021, **60**, 7777–7785.

38 G. Krauss, F. Meichsner, A. Hochgesang, J. Mohanraj, S. Salehi, P. Schmode and M. Thelakkat, *Adv. Funct. Mater.*, 2021, 2010048, DOI: 10.1002/adfm.202010048.

39 H. Jia, Z. Huang, P. Li, S. Zhang, Y. Wang, J. Wang, X. Gu and T. Lei, *J. Mater. Chem. C*, 2021, **9**, 4927–4934.

40 N. Wang, L. Xie, H. Ling, V. Piradi, L. Li, X. Wang, X. Zhu and F. Yan, *J. Mater. Chem. C*, 2021, **9**, 4260–4266.

41 J. J. Samuel, A. Garudapalli, A. A. Mohapatra, C. Gangadharappa, S. Patil and N. P. B. Aetukuri, *Adv. Funct. Mater.*, 2021, 2102903.

42 H. Chen, Y. Guo, G. Yu, Y. Zhao, J. Zhang, D. Gao, H. Liu and Y. Liu, *Adv. Mater.*, 2012, **24**, 4618–4622.

43 K. Shi, W. Zhang, D. Gao, S. Zhang, Z. Lin, Y. Zou, L. Wang and G. Yu, *Adv. Mater.*, 2018, **30**, 1705286–1705293.

44 H. Huang, Z. Chen, R. P. Ortiz, C. Newman, H. Usta, S. Lou, J. Youn, Y.-Y. Noh, K.-J. Baeg, L. X. Chen, A. Facchetti and T. Marks, *J. Am. Chem. Soc.*, 2012, **134**(26), 10966–10973.

45 J. D. Yuen, J. Fan, J. Seifter, B. Lim, R. Hufschmid, A. J. Heeger and F. Wudl, *J. Am. Chem. Soc.*, 2011, **133**, 20799–20807.

46 J. Miao, J. Wang, B. Meng, J. Liu and L. Wang, *Mater. Chem. Front.*, 2018, **2**, 2333–2339.

47 A. Giovannitti, C. B. Nielsen, D. Sbircea, S. Inal, M. Donahue, M. R. Niazi, D. A. Hanifi, A. Amassian, G. G. Malliaras, J. Rivnay and I. McCulloch, *Nat. Commun.*, 2016, **7**, 1–10.

48 E. Wang, Z. Ma, Z. Zhang, K. Vandewal, P. Henriksson, O. Inganäs, F. Zhang and M. R. Andersson, *J. Am. Chem. Soc.*, 2011, **133**, 14244–14247.

49 X. Wu, Q. Liu, A. Surendran, S. E. Bottle, P. Sonar and W. L. Leong, *Adv. Electron. Mater.*, 2020, 2000701.

50 L. Q. Flagg, C. G. Bischak, J. W. Onorato, R. B. Rashid, C. K. Luscombe and D. S. Ginger, *J. Am. Chem. Soc.*, 2019, **141**(10), 4345–4354.

51 A. Savva, R. Hallani, C. Cendra, J. Surgailis, T. C. Hidalgo, S. Wustoni, R. Sheelamanthula, X. Chen, M. Kirkus, A. Giovannitti, A. Salleo, I. McCulloch and S. Inal, *Adv. Funct. Mater.*, 2020, 1907657.

