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# Dielectric Constant Engineering of Organic Semiconductors: Effect of Planarity and Conjugation Length

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## Keywords:

Solar cells, organic semiconductors, homojunction, dielectric constant, molecular geometry.

## Abstract

Bulk heterojunction organic solar cells continue to show steady photoconversion efficiency improvements. However, single component organic solar cells are a particularly attractive alternative due to the relative simplicity of device manufacture. It has been proposed that organic semiconductors with a high dielectric constant ( $\approx 10$ ) could give rise to spontaneous free charge carrier generation upon photoexcitation. In this manuscript we have explored factors that affect the dielectric constant of organic semiconductors, particularly the optical-frequency dielectric constant. We compare the properties of monomers, dimers and trimers of two isoelectronic families of materials that have acceptor units composed of one or two dicyanovinylbenzothiadiazole moieties and one to three donor units. The donor components are composed of either fluorenyl or dithienocyclopentadiene moieties with the same glycol-based solubilising groups. We find that chromophore planarity and orientation with respect to the substrate, and film density affect the optical and electronic properties of the materials, especially the high-frequency dielectric constant. The results also indicate that delocalisation of the highest occupied and lowest unoccupied molecular orbitals is a critical factor. The dimer with two dicyanovinylbenzothiadiazole moieties and two dithienocyclopentadiene units was found to have the highest optical frequency dielectric constant and overall performance.

## Introduction

The bulk heterojunction (BHJ) photoactive layers used in the most efficient organic solar cells are generally composed of at least one electron-donating material and one electron accepting material. The BHJ approach is used to overcome the exciton binding energy and maximise the efficiency of free-charge carrier generation.

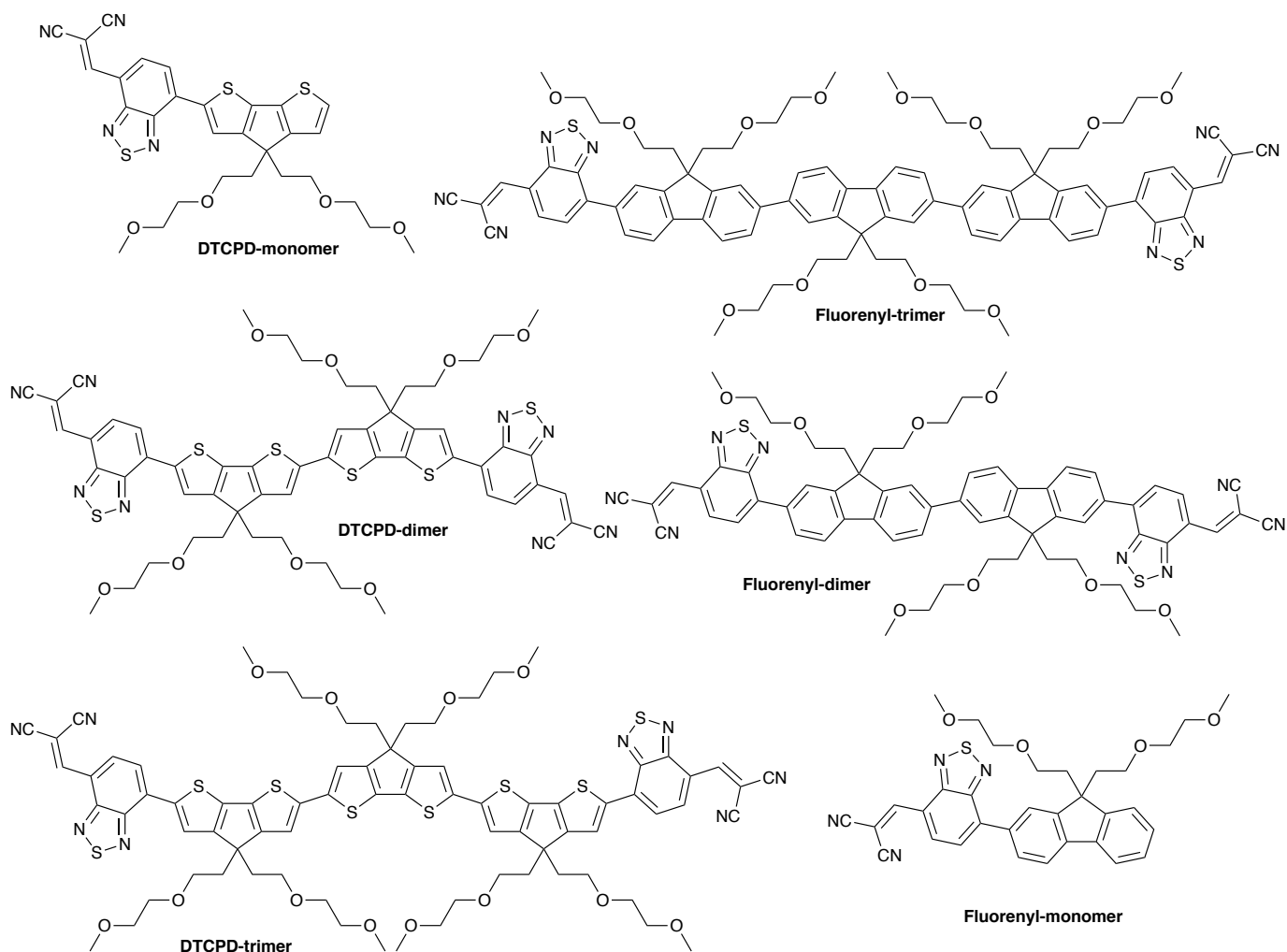
In the past decade, the power conversion efficiency (*PCE*) of solution processed single active layer BHJ solar cells has steadily improved, reaching up to 18%.<sup>[1]</sup> The improvement in BHJ solar cell performance has arisen through a combination of materials design, particularly the development of non-fullerene acceptors,<sup>[2-4]</sup> as well as improvements in device architecture engineering<sup>[5]</sup> and post-processing methods.<sup>[2]</sup> Key to the BHJ solar cell efficiency is control of the active-layer film morphology,<sup>[6-7]</sup> with a delicate balance between the donor and acceptor interfacial area (required for exciton separation) and connected pathways for the free charges to be extracted from the device. However, the performance of a BHJ is sensitive to variations in morphology, which leads to issues with regard to reproducibility in terms of efficiency and device lifetime,<sup>[8]</sup> as well as providing challenges for device scaling, that is, when moving from laboratory testing to large-scale manufacturing. Therefore, in spite of the tremendous efficiencies reported for BHJ solar cells there would be a distinct longer-term advantage of designing materials that could be used in single-component organic solar cells (SCOSCs), that is, solar cells that have a single material in the light-absorbing layer.

The two current approaches to SCOSCs include materials composed of linked donor and acceptor chromophores (an intramolecular “bulk heterojunction”) and materials that have been designed to have a high dielectric constant. Of these two approaches the former is the most advanced in terms of examples of materials and device performance.<sup>[9-12]</sup> The structures of the linked donor and acceptor materials have predominantly had either oligomeric donors connected to fullerene acceptors<sup>[9-10]</sup> or donor polymer cables with perylene diamine acceptor chromophores.<sup>[11-12]</sup> The latter approach has led to a strategy of morphology control through thermally driven self-organised phase separation, with good device stability under long-term illumination being demonstrated.<sup>[12]</sup>

The alternative less developed approach to homojunction devices is to design organic semiconductor materials that have a high dielectric constant ( $\epsilon$ ). Seminal papers proposing this approach reported that increasing the dielectric constant of the light-absorbing organic semiconductor to around 10 could lower the exciton binding energy ( $E_b \propto \frac{1}{\epsilon^2}$ ) such that free charge carriers would be generated upon photoexcitation in a similar manner to inorganic and hybrid semiconductor materials.<sup>[13-14]</sup> However, the dielectric constant of a material is frequency dependent and a key question to consider in the development of such materials is which frequency or frequencies are critical for charge generation and/or extraction. An early and commonly reported strategy to increase the dielectric constant has been to replace the alkyl groups normally used to solubilise organic semiconductors with ethylene glycol moieties.<sup>[15-17]</sup> This approach was particularly successful in increasing the  $\epsilon$  at low-frequency ( $\epsilon_{lf}$ ) due to enhanced dipole polarisability, with an  $\epsilon_{lf}$  as high as 9.8 being reported for a small molecule.<sup>[15]</sup> An important outcome of the early work on increasing the low-frequency dielectric constant was that the ethylene glycol moieties did not change the fundamental spectral, electronic or charge-transporting properties of the materials.<sup>[15, 17-18]</sup> Subsequent work reported that the optical frequency dielectric constant ( $\epsilon_{opt}$ ) was important for the direct generation of free charge carriers

upon photoexcitation.<sup>[19-20]</sup> There are a number of factors that can affect the optical frequency dielectric constant, including chromophore conjugation length and orientation (with respect to the substrate), the dipole moment of the material, and film density.<sup>[21]</sup> There has been one report showing the effect of conjugation length, whereby the optical frequency dielectric constant of a dimeric material composed of two dithienocyclopentadiene “donor” and two dicyanovinylbenzothiadiazole “acceptor” units was found to be higher than the corresponding monomer material (one of each). The optical frequency dielectric constant was found to increase from 3.6 to 4.6, and interestingly, homojunction solar cells showed that charges were generated more efficiently near the optical gap for the dimer.<sup>[19]</sup> That is, the results imply that extending the conjugation length can lead to an increased optical frequency dielectric constant and more efficient charge generation. However, the *PCEs* of these homojunction devices have yet to be reported. A further outcome of the early study was that solar cells composed of the glycolated equivalents of the monomer and dimer materials had higher external quantum efficiencies near the absorption onset compared to their alkylated equivalents. One potential explanation for this latter result is that the flexibility of the glycol side chains leads to less steric hindrance,<sup>[18, 22]</sup> resulting in closer  $\pi$ - $\pi$  stacking and thus a higher film density.<sup>[19]</sup>

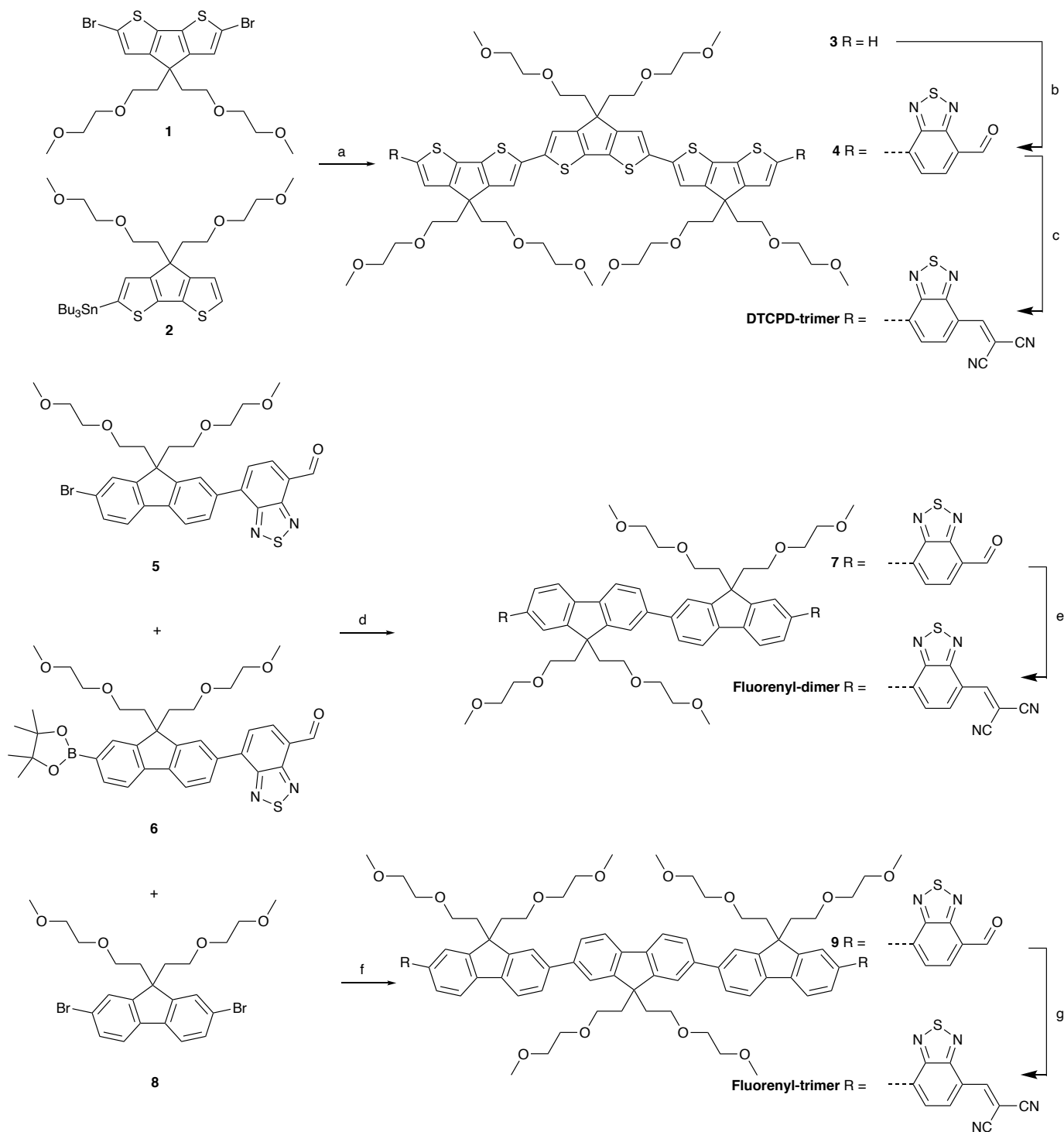
In this manuscript we comprehensively explore the effect of the chromophore conjugation length and planarity on the optoelectronic properties of two families of isoelectronic semiconductor materials. The materials have acceptor units composed of one or two dicyanovinylbenzothiadiazole moieties and one to three donor units. The donor components are composed of either fluorenyl or dithienocyclopentadiene moieties with the same glycol-based solubilising groups **Figure 1**. We report the dielectric constants of the materials and how these relate to their chemical and electronic structure, and film morphology. Finally, we show how the planarity, and chromophore conjugated length and type affects the charge transport properties of the materials and their performance in homojunction solar cells.



**Figure 1.** Structures of the compounds compared in this study.

## Results and discussion

### Synthesis



**Scheme 1.** Synthesis of DTCPD-trimer, and Fluorenyl-dimer and -trimer. *Reagents and conditions:* a)  $\text{Pd}(\text{PPh}_3)_2\text{Cl}_2$ , toluene, reflux, 21 h; b) 1) *n*-butyllithium, tetrahydrofuran,  $-78\text{ }^\circ\text{C}$ , 1 h, then, tri-*n*-butyltin chloride, room temperature, 1.5 h, then  $\text{Pd}(\text{PPh}_3)_4$ , toluene, reflux, 16 h; c) malononitrile, pyridine, toluene,  $60\text{ }^\circ\text{C}$ , 5 h; d)  $\text{Pd}(\text{PPh}_3)_4$ ,  $\text{K}_2\text{CO}_3$  (2M aq.), toluene, *t*-BuOH, reflux, overnight; e) malononitrile, pyridine, chlorobenzene,  $75\text{ }^\circ\text{C}$ , overnight; f)  $\text{Pd}(\text{PPh}_3)_4$ ,  $\text{Cs}_2\text{CO}_3$ , toluene, reflux, 22 h; g) malononitrile, pyridine, chloroform, reflux, 3.5 h.

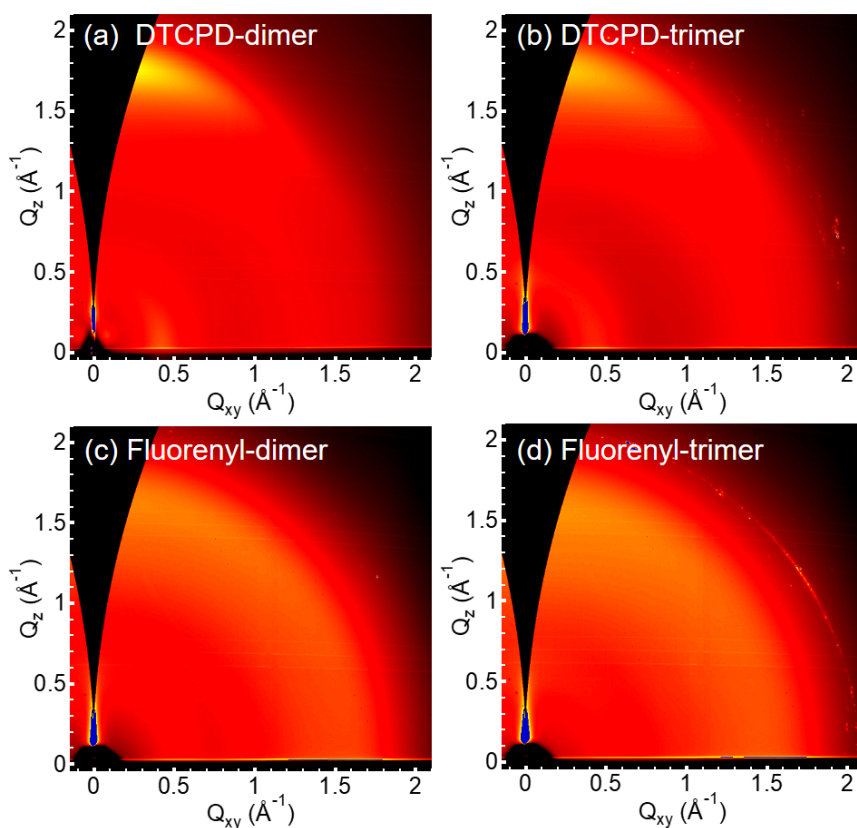
The syntheses of the **DTCPD-monomer**<sup>[15]</sup> and **DTCPD-dimer**<sup>[19]</sup> and **Fluorenyl-monomer**<sup>[15]</sup> have been previously reported. The synthetic pathways to the **DTCPD-trimer** and **Fluorenyl-dimer** and **-trimer** from reported intermediates are shown in **Scheme 1**. The first step in the synthesis of the **DTCPD-trimer** was the

Stille coupling of **1**<sup>[19]</sup> and **2**<sup>[15]</sup> to form trimer **3**, which was achieved a 20% yield. Trimer **3** was converted to the bis-stannyl derivative and then reacted under palladium-catalysed Stille conditions with 7-bromobenzo[*c*][1,2,5]thiadiazole-4-carbaldehyde to give **4** in a 40% yield. The final step required the condensation of **4** with malononitrile, with the **DTCPD-trimer** being formed in an isolated yield of 56%. The syntheses of the **Fluorenyl-dimer** and **-trimer** utilised a similar strategy, with **5** being the key intermediate. **5** was formed in an 52% yield through a chemoselective Suzuki reaction between 2-bromo-7-iodo-9,9-bis[2-(2-methoxyethoxy)ethyl]-9*H*-fluorene (see Supporting Information) and 7-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)benzo[*c*][1,2,5]thiadiazole-4-carbaldehyde.<sup>[23]</sup> To prepare the **Fluorenyl-dimer** and **-trimer**, **5** was converted into the corresponding boronate ester **6**, which due its instability was then used immediately. In the case of the **Fluorenyl-dimer**, **5** was reacted with **6** to form dialdehyde **7** (59% yield) and then in the final step **7** underwent a Knoevenagel condensation with malononitrile, giving the **Fluorenyl-dimer** in a 37% yield. For the synthesis of the **Fluorenyl-trimer**, **6** was reacted with the glycolated dibromofluorene derivative **8** to give dialdehyde **9** in a 35% yield. Finally, **9** was condensed with malononitrile to give the **Fluorenyl-trimer** in a 62% yield.

### Film density and morphology

To set the framework for the differences in the optoelectronic properties of the materials we first determined how the materials packed in films using X-ray reflectometry (XRR) and grazing-incidence wide angle X-ray scattering (GIWAXS) measurements to determine the density and  $\pi$ -stacking distances. The density of the DTCPD-based monomer, dimer and trimer from XRR measurements were 1.23 g/cm<sup>3</sup>, 1.29 g/cm<sup>3</sup> and 1.29 g/cm<sup>3</sup>, respectively. Apart from the **Fluorenyl-monomer** (1.23 g/cm<sup>3</sup>), the density of the **Fluorenyl-dimer** and **Fluorenyl-trimer** (1.13 g/cm<sup>3</sup>) were lower than their DTCPD counterparts. The scattering patterns from the GIWAXS measurements of the dimers and trimers of both families are shown in **Figure 2**. The **DTCPD-dimer** and **-trimer** exhibit reflections that are indexed to lamellar stacking (100) and  $\pi$ - $\pi$  stacking (010) peaks at  $\sim 0.42 \text{ \AA}^{-1}$  and  $1.8 \text{ \AA}^{-1}$ , respectively. The appearance of the (100) peaks in-plane and (010) peaks out-of-plane indicate that the DTCPD-based materials adopt a face-on orientation with respect to the substrate. The appearance of the (100) peak at  $\sim 0.42 \text{ \AA}^{-1}$  corresponds to a lamellar (side-chain) stacking repeat distance of  $\sim 15 \text{ \AA}$ , while the appearance of the (010) peak at  $1.77 \text{ \AA}^{-1}$  corresponds to a  $\pi$ - $\pi$  stacking distance of  $3.55 \text{ \AA}$  for both DTCPD-based molecules. The (100) and (010) peaks for the **DTCPD-dimer** are both narrower than those of the **DTCPD-trimer** indicating that the dimer (see **Figure S1** in the Supporting Information) has improved crystallinity (either larger crystallites and/or reduced crystalline disorder). In contrast, the fluorenyl-based materials are essentially amorphous, indicated by the lack of well-defined scattering features. A very broad ring centred at  $\sim 1.6 \text{ \AA}^{-1}$  is seen, characteristic of weakly correlated  $\pi$ - $\pi$  stacking corresponding to an average  $\pi$ - $\pi$  stacking distance of  $4.0 \text{ \AA}$ , which is consistent with the lower film density. The fluorenyl-based materials also exhibit a lack of texture indicating a lack of preferential molecular orientation in thin film. Based on the structural analysis and the criteria postulated to affect the optical-frequency dielectric constant [chromophore conjugation length and orientation (with respect to the

substrate), and film density]<sup>[21]</sup> it might be expected that the DTCPD-based materials would outperform the fluorenyl-containing compounds across a range of the critical optoelectronic metrics.



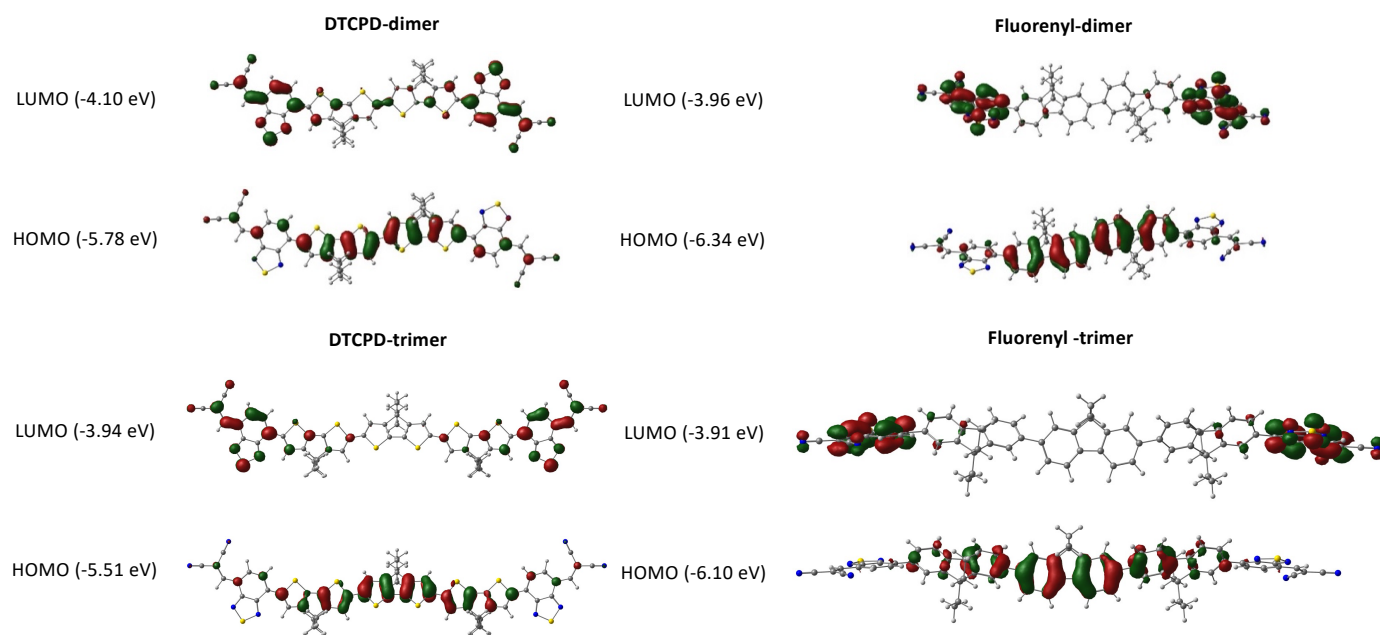
**Figure 2.** 2D GIWAXS patterns of neat films of **DTCPD-dimer** (a) and **-trimer** (b) and **Fluorenyl-dimer** (c) and **-trimer** (d) cast from chloroform solutions onto silicon substrates.

### Molecular orbital calculations

To gain insight into the differences in densities and  $\pi$ -stacking distances between the different materials classes, we undertook density functional theory (DFT) calculations. The starting structure in each case had the molecule in a planar arrangement, which was then allowed to relax to its lowest energy ground-state geometry using the B3LYP functional and DZP Slater-type basis set in the gas phase, with the results for the dimers and trimers shown in **Figure 3**. For computational efficiency the glycol units were replaced by methyl groups. The key difference between the DTCPD- and fluorenyl-based materials was that the former had greater planarity overall. That is, the dihedral angle between the benzothiadiazole and DTCPD unit was  $0.04^\circ$  (compared to  $38^\circ$  for the fluorenyl-benzothiadiazole unit) and the dihedral angle between the DTCPD units was only  $18^\circ$  compared to  $44^\circ$  for the **Fluorenyl-dimer**. The results are logical in that the steric interactions of the protons that lead to the twisting between the adjacent units would be expected to be less for the DTCPD moieties than the fluorenyl units. The fact that the **DTCPD-dimer** and **-trimer** are more planar than their fluorenyl counterparts is also consistent with the measured higher film density and smaller  $\pi$ -stacking distance for the former. A further consequence of the difference in the planarity of the two families of materials is the distribution of the highest occupied (HOMO) and lowest unoccupied molecular orbitals



(LUMO) in the ground-state structures shown in **Figure 3**. For the **DTCPD-dimer** there is HOMO and LUMO density on both the DTCPD and benzothiadiazole moieties. In contrast, the **Fluorenyl-dimer** has the HOMO and LUMO density strongly localised on the fluorenyl and benzothiadiazole units, respectively. For the **DTCPD-trimer** the HOMO is delocalised over all five units while the LUMO has no density on the central DTCPD-unit. In the case of the **Fluorenyl-trimer**, the HOMO had greatest density on the central fluorenyl moiety and little delocalisation of the LUMO across any of the fluorenyl units. Finally, the HOMO–LUMO gap was found to decrease for both families in moving from the dimer to the trimer.

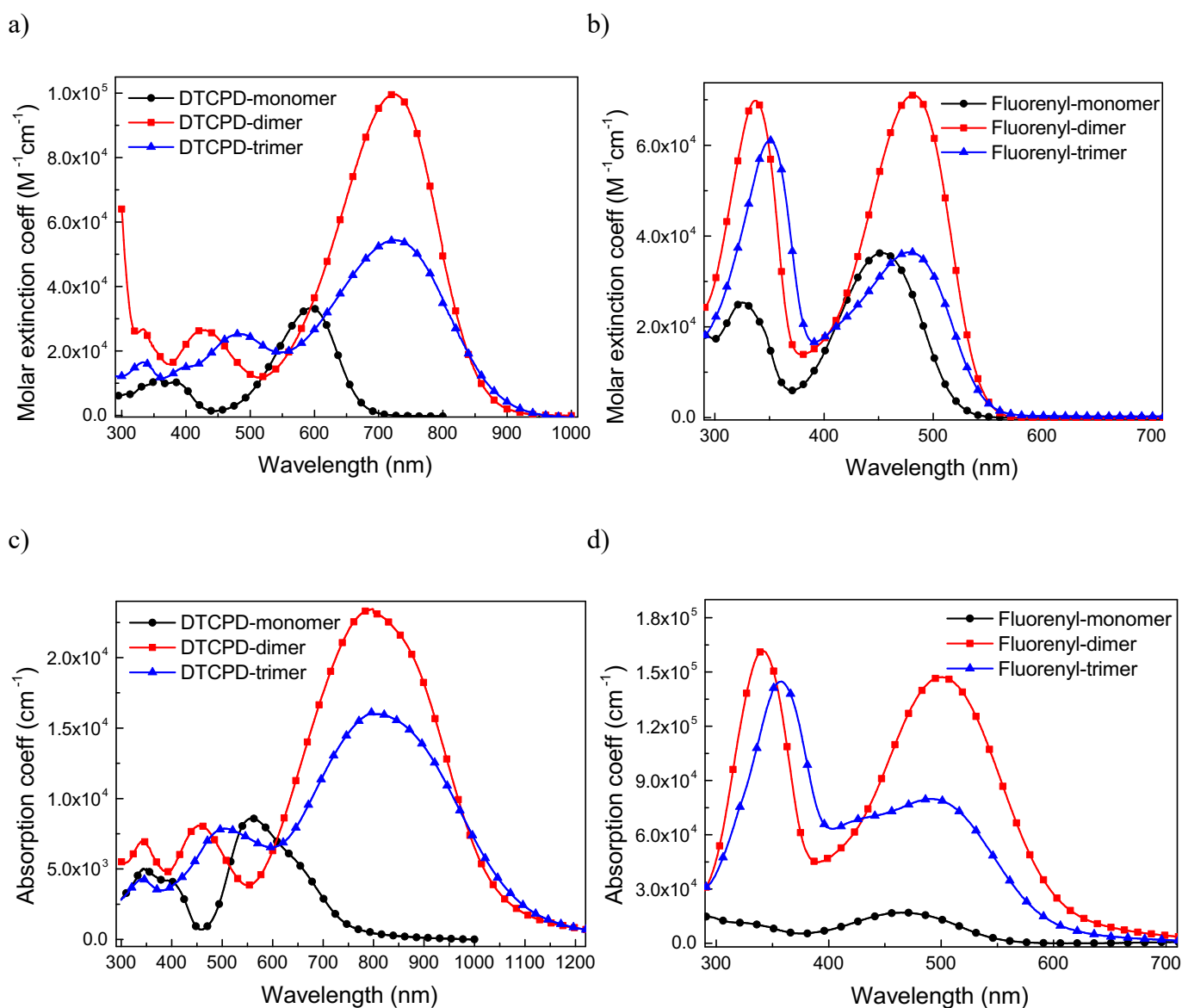


**Figure 3.** Molecular orbitals of the DTCPD- and fluorenyl-based dimers and trimers showing the HOMO and LUMO distributions. Calculations were done using the Amsterdam Density Functional package (ADF 2019.301).

### Optical properties

In the next part of the analysis of the properties of the materials we compared the effect of extending the conjugation length of the two families on the absorption spectra (**Figure 4**). The solution absorption spectra of the DTCPD series are shown in **Figure 4a** and it can be seen that in moving from the monomer to the dimer there is a significant red shift in the onset and peak of the long-wavelength absorption ( $\lambda_{\text{max}}$  from 595 nm to 724 nm), with a negligible change in moving to the trimer ( $\lambda_{\text{max}} = 726$  nm). The fact that the long wavelength  $\lambda_{\text{max}}$  was not red shifted further for the trimer and its smaller molar extinction coefficient than the dimer is consistent with the molecular orbital calculations that showed there was no LUMO density on the central DTCPD unit. A similar trend was observed for the fluorenyl series – there was a significant red shift in the long-wavelength absorption peak in moving from the monomer to the dimer ( $\lambda_{\text{max}} = 451$  nm to 482 nm), whereas the  $\lambda_{\text{max}}$  for the trimer was at a similar wavelength to the dimer ( $\lambda_{\text{max}} = 479$  nm) (**Figure 4b**). The onset of the absorption of the DTCPD-based materials being at longer wavelengths than those of

the fluorenyl-containing materials is consistent with the relative energy levels of the individual donor units (DTCPD *versus* fluorene) and the HOMO and LUMO energies from the DFT calculations.



**Figure 4.** Molar extinction coefficient as a function of wavelength for solutions of the DTCPD (a) and fluorenyl (b) series, and absorption coefficient as a function of wavelength for films of the DTCPD (c) and fluorenyl (d) series.

The behaviour of the absorption spectra of the two families were completely different when moving to the solid state. In the case of the fluorenyl series there was a small red shift in the onset of the absorption and long-wavelength peak, which is normally ascribed to slight planarisation of the chromophores in the film (**Figure 4d**). In contrast, for the DTCPD family of materials there was a significant red shift in the absorption onset ( $>100$  nm) for the dimer and trimer, with the film absorption in both cases going beyond 1100 nm (1.1 eV) (**Figure 4c**). Planarisation of the chromophore alone would not be expected to give rise to such a large red shift and hence the observed changes in the absorption spectra of the DTCPD-based materials are likely to be due to a combination of planarisation and their propensity to  $\pi$ -stack, as observed from the X-ray scattering experiments.

## Electronic properties

The redox properties of the materials were determined in solution using cyclic voltammetry, with the  $E_{1/2s}$  reported against the ferrocene/ferrocenium couple and the results summarised in **Table 1**. The DTCPD-based materials were found to undergo chemically reversible oxidations (one, two and three, respectively) and two chemically reversible reductions. In contrast, only the **Fluorenyl-monomer** underwent a chemically reversible oxidation, with the monomer, dimer and trimer undergoing one, two and two chemically reversible reductions, respectively. The first oxidation (for the monomer) and reduction potentials of the fluorenyl-based materials were at more positive and negative potentials compared to those of the DTCPD containing materials, which is consistent with the trend in the calculated HOMO and LUMO energies. The thin film ionisation potentials (IPs) of the materials were also measured using photoelectron spectroscopy in air (PESA) (**Table 1**). The IPs of the fluorenyl-based materials we found to be larger (5.7-5.9 eV) than those of the compounds composed of the DTCPD units (5.0-5.3 eV).

**Table 1** Summary of the optical and electronic properties of the materials.

Materials	DTCPD-			Fluorenyl-		
	monomer	dimer	trimer	monomer	dimer	trimer
$\lambda_{\max}$ (solution/film, nm)	595/612	724/799	726/799	451/469	482/501	479/492
IP (eV)	5.3	5.0	5.0	5.9	5.9	5.7
$E_{1/2(\text{ox})}$ (V)	0.5	0.2, 0.9	0.05, 0.2, 0.9	0.6	–	–
$E_{1/2(\text{red})}$ (V)	-1.2	-1.1, -1.5	-1.1, -1.7	-1.3	-1.2, -1.8	-1.2, -1.8, -2.1
Optical gap (eV)	1.9	1.2	1.1	2.4	2.2	1.9

The dielectric constants of the materials were then determined using charge extraction with linearly increasing voltage (CELIV) and impedance measurements for  $\epsilon_{\text{if}}$ , and a combination of spectroscopic reflectometry, transmission and ellipsometry for  $\epsilon_{\text{opt}}$ , with the results shown in **Table 2**. The optical-frequency dielectric constant was higher for all the DTCPD materials compared to the equivalent fluorenyl-based materials, with the **DTCPD-dimer** and **-trimer** having the highest  $\epsilon_{\text{opt}}$  of 4.6. At this point it is worth remembering the characteristics that are proposed to be critical for a molecule to have a high optical frequency dielectric constant, namely, chromophore conjugation length ( $\pi$ -delocalisation) and orientation (with respect to the substrate), and film density.<sup>[21]</sup> Based on these criteria it is not surprising that the fluorenyl family of materials have the smaller optical frequency dielectric constants. The X-ray scattering

data show that the **Fluorenyl-dimer** and **-trimer** have the lower density and larger  $\pi$ - $\pi$  stacking distance when compared to their DTCPD counterparts. Furthermore, the GIWAX data show that the films composed of the **Fluorenyl-dimer** and **-trimer** are amorphous with no preferential orientation of the chromophore with respect to the substrate. In contrast, the **DTCPD-dimer** and **-trimer** have a preferred face-on orientation with respect to the substrate. Perhaps the more surprising result is that the optical frequency dielectric constant of the **DTCPD-trimer** is not larger than the dimer in spite of its larger conjugated framework. In the case of the **DTCPD-dimer** the HOMO and LUMO density is distributed across all the sub-units. However, while the HOMO of the **DTCPD-trimer** is delocalised over the trimer chromophore, the LUMO is not, with the central ring bereft of LUMO density. These results suggest that to increase the dielectric constant both the HOMO and LUMO need to be as fully delocalised as possible.

Finally, before fabricating homojunction solar cells we measured the hole and electron mobilities of the dimers and trimers using metal-insulator-semiconductor charge extraction by linear increasing voltage (MIS-CELIV), with the results shown in **Table 2** and **Figure S2**. The two monomers were previously reported to have similar electron mobilities ( $\approx 10^{-6}$  cm<sup>2</sup>/Vs) but did not have a measurable hole mobility.<sup>[15]</sup> In fact, using MIS-CELIV we did not observe any hole mobility for the fluorenyl-based materials. Interestingly, for the fluorenyl-series, increasing the conjugation length of the chromophore led to a decrease in the electron mobility. Presumably this arises from the fact that the LUMO density is strongly localised on the end dicyanovinylbenzothiadiazole units of the molecules leading to decreased probability of intermolecular overlap of these units as they become a smaller component of the overall molecule (i.e., there are larger numbers of fluorenyl units). In contrast, both the **DTCPD-dimer** and **-trimer** have measurable hole mobilities of order  $\approx 10^{-5}$  cm<sup>2</sup>/Vs, with the dimer having an order of magnitude higher electron mobility at  $1 \times 10^{-4}$  cm<sup>2</sup>/Vs. The electron mobility of the **DTCPD-trimer** is over an order of magnitude lower than the dimer at  $6.5 \times 10^{-6}$  cm<sup>2</sup>/Vs, which would be consistent with the more localised nature of the LUMO of the former, decreasing the probability of intermolecular LUMO overlap in the thin film.

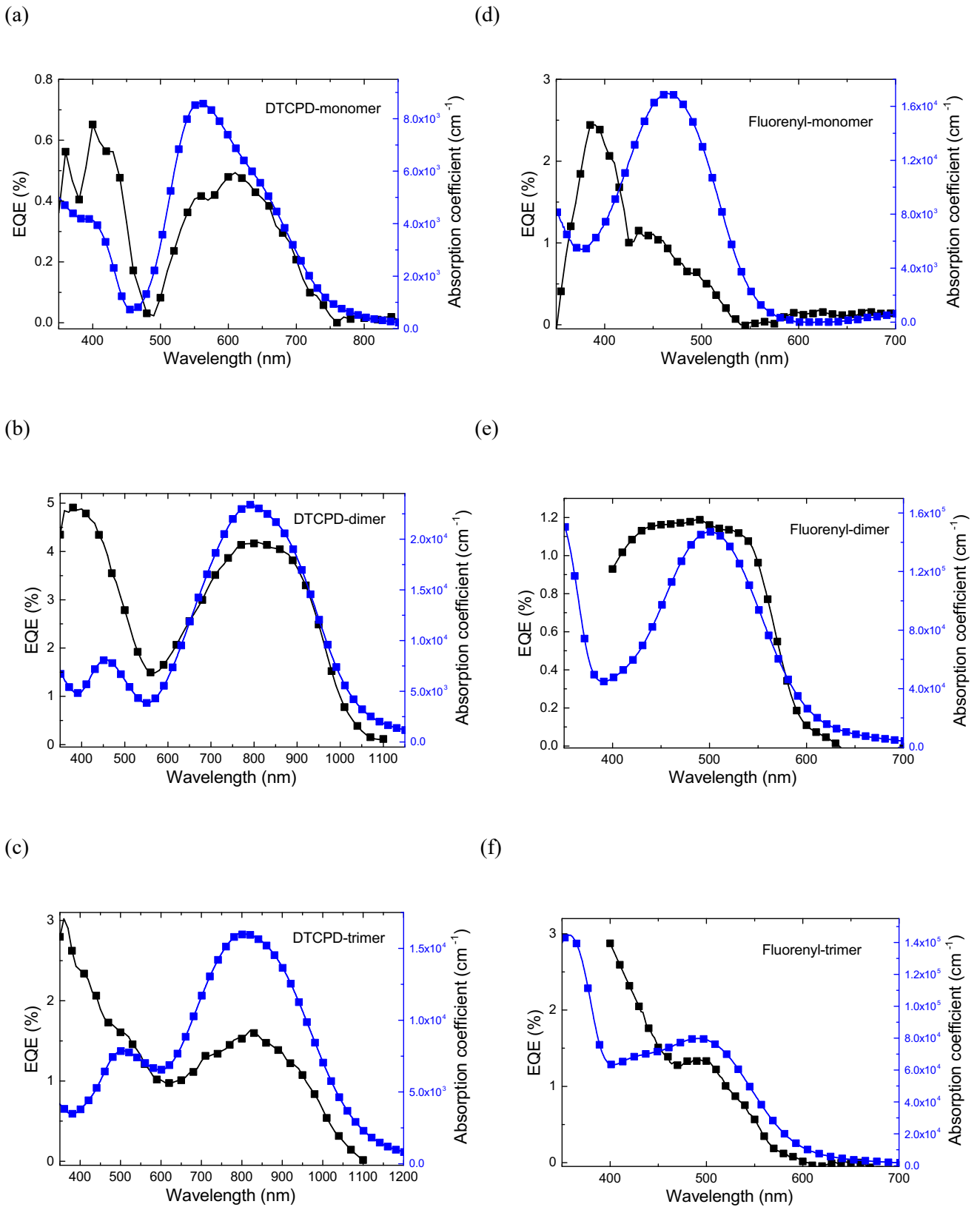
**Table 2** Summary of the electronic properties, diode mobility, film density,  $\pi$ -stacking distance and homojunction device parameters of the compounds for comparison in this study.

Materials	DTCPD-			Fluorenyl-		
	monomer	dimer	trimer	monomer	dimer	trimer
$\epsilon_{lf}$	9.8	6.1	5.1	8.5	3.5	5.0
$\epsilon_{opt}$	3.2	4.6	4.6	2.9	3.0	3.0
$\mu_h$ (cm <sup>2</sup> /Vs)	–	$1.25 \times 10^{-5}$	$1.5 \times 10^{-5}$	–	–	–
$\mu_e$ (cm <sup>2</sup> /Vs)	$3.0 \times 10^{-6}$	$1.0 \times 10^{-4}$	$6.5 \times 10^{-6}$	$1.3 \times 10^{-6}$	$1.4 \times 10^{-8}$	$4.2 \times 10^{-7}$
Density (g/cm <sup>3</sup> )	1.23	1.29	1.29	1.23	1.15	1.13

$\pi$ -stacking distance (Å)	3.88	3.59	3.58	3.92	3.83	4.05
EQE (%) at the absorption peak	0.5	4.2	1.6	0.8	1.2	1.3
$J_{sc}$ (mA/cm <sup>2</sup> )	0.06±0.00	1.11±0.03	0.57±0.02	0.02±0.00	0.15±0.01	0.17±0.01
$V_{oc}$ (V)	0.32±0.06	0.56±0.02	0.31±0.03	0.29±0.06	0.58±0.04	0.71±0.10
$FF$ (%)	30.4±1.4	34.9±0.6	33.6±0.4	32.4±2.3	25.0±0.7	23.2±1.2
$PCE$ (%)	0.006±0.002	0.21±0.01	0.06±0.01	$2 \times 10^{-3} \pm 0.00$	0.02±0.00	0.03±0.00
Hero $PCE$ (%)	0.008	0.23	0.06	0.002	0.025	0.03

## Devices

Based on the optical and electronic properties it would be expected that DTCPD-based materials would provide better performance in homojunction solar cells. The DTCPD-materials absorb across a broader wavelength range, have higher optical frequency dielectric constants, and in the case of the **DTCPD-dimer** and **-trimer** are capable of both hole and electron transport. Homojunction devices with structure ITO/PEDOT:PSS/homojunction/Ca/Al were fabricated and tested, with the EQEs *versus* absorption spectra shown in **Figure 5a-5f**, current density versus voltage ( $J$ - $V$ ) in **Figure S3**, and a summary of the device characteristics in **Table 2**. A key feature of all the devices was that independent of the  $\epsilon_{opt}$  they all generated charge near the optical gap, with those of the **DTCPD-dimer** and **-trimer** having the highest EQE near the optical gap. The devices comprising the monomers showed diode characteristics but low performance, which can at least in part be ascribed to the lack of hole mobility and poor electron mobility. **The DTCPD-dimer** showed good diode characteristics and low dark current leakage. In contrast, the **Fluorenyl-dimer** (and indeed **-trimer**) had very poor performance, but they also had the lowest electron mobilities and no measurable hole mobility. The devices containing the **DTCPD-trimer** were found to have higher current leakage than those incorporating the dimer, which affected the fill factor ( $FF$ ). Nevertheless, these results suggest that there might be a minimal  $\epsilon_{opt}$  to achieve charge generation near the optical gap, with the higher the  $\epsilon_{opt}$  the more efficient the process. The homojunction device containing the **DTCPD-dimer** was found to have the highest photoconversion efficiency ( $PCE$ ) of 0.23%. While this is low compared to the state-of-the-art intramolecular BHJ SCOSCs, it is comparable to the very early reports of non-fullerene acceptor-based devices.<sup>[24-26]</sup> The **DTCPD-dimer** devices had the largest  $J_{sc}$  (consistent with the relative high film absorption coefficient and onset >1100 nm) and  $FF$  (consistent with the higher and more balanced electron and hole mobilities). A clear trend was that the more disordered and poorly aligned a material was (relative to the substrate) the poorer the device performance.



**Figure 5.** Film absorption spectra and external quantum efficiency (EQE) of homojunction devices with a structure of ITO/PEDOT:PSS/homojunction/Ca/Al composed of **DTCPD-monomer** (a), **-dimer** (b) and **-trimer** (c), and **Fluorenyl-monomer** (d), **-dimer** (e) or **-trimer** (f).

## Conclusions

We have developed two families of materials designed to provide a framework and insight into the creation of high optical frequency dielectric constant materials that could be used in simple homojunction organic solar cells. While a number of factors that can affect the optical frequency dielectric constant have been proposed, including chromophore conjugation length and orientation (with respect to the substrate), and film density, there has not been a sufficient body of evidence as to which of these are most important. Our results show that the planarity of the chromophore is critical to achieving the desired properties. Planarity increases molecular orbital delocalisation, the film density and the possibility of favourable orientation of the chromophore with respect to the substrate, leading to increased light absorption, film crystallinity, and charge mobility. However, the results show that increasing the conjugation length does not always lead to an increase in dielectric constant or an improvement in the optoelectronic properties, which appears to be directly related to the degree of molecular orbital delocalisation. The dimer with two dicyanovinylbenzothiadiazole moieties and two dithienocyclopentadiene was found to have the highest optical-frequency dielectric constant, relatively balanced and reasonable hole and electron mobilities and best overall homojunction device performance. While the photoconversion efficiency of the device was significantly lower than current state-of-the-art organic solar cells it is interesting note that early non-fullerene-acceptor-based devices had similar performance. In conclusion, the results from this work add extra key criteria that need to be considered when developing high dielectric constant materials for homojunction organic solar cells.

## Experimental

**Material Synthesis and Characterisation.** Commercial materials were used as received unless otherwise stated. Dichloromethane used for the optical measurements was dried over calcium hydride or potassium carbonate and distilled immediately prior to use. Dichloromethane (HPLC grade,  $\geq 99.9\%$ ) and methanol (HPLC grade,  $\geq 99.9\%$ ) used for the mass spectroscopy measurements were used as received from Sigma Aldrich<sup>®</sup>. Dichloromethane for column chromatography was distilled by rotary evaporation under reduced pressure before use. Column chromatography was performed with Merck silica, 230-400 mesh. When solvent mixtures are used, the proportions are given by volume. Medium pressure liquid chromatography (MPLC) was performed on a Grace Reveleris X2 system using silica or C18 reverse-phase cartridges. Size-exclusion column chromatography was performed with Bio-Rad Bio-Beads S-X1, 200-400 mesh.

<sup>1</sup>H and <sup>13</sup>C NMR spectra were recorded on Bruker AS500, AV500 or AN300 spectrometers. Chemical shifts are reported in parts per million (ppm) and are referenced to the residual solvent peak (CDCl<sub>3</sub> 7.26 ppm for <sup>1</sup>H or 77.0 ppm for <sup>13</sup>C, and CD<sub>2</sub>Cl<sub>2</sub> 5.32 ppm for <sup>1</sup>H or 53.8 ppm for <sup>13</sup>C). GI-H = H on glycol solubilising groups, DTCPD-H = dithienocyclopentadienyl H, FI-H = fluorenyl H, BT-H = benzothiodiazolyl H, and VIN-H = vinyl H. Multiplicities are reported as singlet (s), doublet (d), doublet of doublets (dd), and multiplet (m) and coupling constants (*J*) are given to the nearest 0.5 Hertz (Hz). High resolution electrospray

ionisation (HRMS) accurate mass measurements were recorded in positive mode on a Bruker MicroTOF-Q (quadrupole time-of-flight) instrument with a Bruker ESI source. Elemental analyses were carried out using a Carlo Erba NCHS Analyser Model NA 1500 instrument at the School of Chemistry and Molecular Biosciences, The University of Queensland. Infrared (IR) absorption spectra were measured on a Perkin-Elmer Spectrum 100 FT-IR spectrometer as neat samples using an ATR crystal. Differential Scanning Calorimetry (DSC) was performed on PerkinElmer® Pyris Diamond DSC. Thermogravimetric analysis (TGA) was carried out on a Perkin-Elmer STA 6000 Simultaneous Thermal Analyser. Decomposition temperatures ( $T_d$ ) are reported for a 5% decrease in mass. Melting points were measured using a Büchi Melting Point B-545 and are uncorrected. Absorption spectra were measured using a Varian Cary 5000 UV-vis-NIR spectrophotometer in double distilled dichloromethane (calcium hydride or potassium carbonate) or spun onto quartz substrates (from chlorobenzene) for solid-state measurements. Absorbance shoulders are denoted as sh. Cyclic voltammetry measurements were carried out on a BASi Epsilon Analyzer at room temperature in an electrolyte solution of 0.1 M tetra-*n*-butylammonium perchlorate in argon-purged anhydrous tetrahydrofuran or dichloromethane. Glassy carbon working, platinum auxiliary, and silver/silver nitrate in acetonitrile reference electrodes were used, and the  $E_{1/2}$  values are quoted relative to the ferrocene/ferrocenium couple. Electrodes were polished between measurements with a polishing pad wetted with deionised water. The ionisation potentials were measured using Photoelectron Spectroscopy in Air (PESA) on a Riken Keiki AC-2 PESA spectrometer with a power setting of 5-10 nW and a power number of 0.5.

**Device Characterisation.** 20 Ohm square<sup>-1</sup> indium tin oxide (ITO) coated glass substrates (Xinyan, Hong Kong) were pre-cleaned using an Alconox (detergent) solution held at 70 °C before being sonicated in sequence with Alconox, de-ionised water, acetone and 2-propanol for 6 min each. The substrates had dimensions of 2.5 cm × 2.5 cm with six 0.2 cm<sup>2</sup> active area pixels. A 30 nm PEDOT:PSS layer was spin-coated onto the cleaned ITO glass followed by baking at 150 °C for 15 mins in the air, and then the active material solutions were spin-coated onto the PEDOT:PSS layer in an MBraun glove box with oxygen concentration < 1 ppm and water concentration < 1 ppm. Solutions of the active materials were prepared in chloroform at a concentration of 10 mg/mL. The solutions were filtered and then diluted to 5 mg/mL with chloroform. The solutions were spin-coated at 1000 rpm for 30 s to give films with a thickness of around 40-50 nm. The films were not thermally annealed after spin-coating and the thicknesses were measured using a Veeco Dektak 150 profilometer. Finally, 15 nm Ca and 100 nm Al were sequentially thermally deposited under a 10<sup>-6</sup> mbar vacuum.

The devices were tested into a nitrogen filled MBraun glove box with oxygen and water concentrations of <1 ppm. Current-voltage ( $J-V$ ) characteristics were acquired using a Keithley 2400 Source Measurement Unit with a 4-wire configuration to eliminate the effect of the cable resistances and SMU internal impedance in the measurement circuit. The simulated Air Mass 1.5 Global (AM 1.5 G) illumination was provided by an Abet ABA Sun 2000 Solar Simulator. The illumination intensity of 1.3 sun used for the organic solar cell



measurements was set using a paired Newport calibrated meter in conjunction with an unfiltered silicon reference cell. Due to higher IR intensity, the 1.3 sun illumination from an unfiltered silicon reference cell was required such that 1 sun illumination corresponding to KG5 Si reference cell could be obtained and the  $J_{SC}$  from the organic solar cells equalled that determined from the integrated EQE. The  $J-V$  curves were measured on at least six devices to provide relevant statistics. An aperture of 0.2 cm<sup>2</sup> was applied during illumination test. The external quantum efficiency (EQE) spectra were recorded with a PV Measurement QEX7 setup, which was calibrated using an NREL certified photodiode and operated without white light bias and chopped and locked in the small perturbation limit.

**Mobility measurements.** The diode mobility for the neat films and blend films were measured using MIS-CELIV. An arbitrary waveform generator (Agilent 33250A) was used to generate the CELIV triangle pulse with adjustable voltage slope, offset and pulse length. The signal was recorded using a digital storage oscilloscope (LeCroy Waverunner A6200). The MIS-CELIV devices were fabricated with a structure of ITO/BCB (30 nm)/MgF<sub>2</sub> (40 nm)/semiconductor layer/Al for electron mobility and ITO/BCB (30 nm)/MgF<sub>2</sub> (40 nm)/semiconductor layer/MoOx/Ag for hole mobility. The thickness of the semiconductor layer for the MIS-CELIV measurements was around 150 nm.

**GIWAXS measurements.** GIWAXS measurements were conducted at the SAXS/WAXS beamline of the Australian Synchrotron.<sup>[27]</sup> Highly collimated 9 keV X-rays were calibrated to be at a tilt angle of  $0 \pm 0.01$  degrees when parallel to the surface of each sample by use of a Silicon crystal analyzer. A Dectris Pilatus 1M detector collected 2D scattering patterns. Each scattering pattern was tiled together from three 1 second images with the detector slightly moved between exposures, such that the resulting image removes gaps between the detector modules. The sample to detector distance was measured using a silver behenate scattering standard. The scattering data reported were acquired at the critical angle identified as the angle with the highest scattering intensity. Data was analysed using a modified version of the NIKA small angle scattering analysis package.<sup>[28]</sup>

## Synthesis.

### **4,4,4',4',4'',4''-Hexakis[2-(2-methoxyethoxy)ethyl]-4*H*,4'*H*,4''*H*-2,2':6',2''-tercyclopenta[2,1-*b*:3,4-*b'*]dithiophene (3)**

A mixture of **1**<sup>[19]</sup> (354 mg, 0.66 mmol), **2** (940 mg, freshly prepared following the procedure described in [15], not purified) and bis(triphenylphosphine)palladium(II) dichloride (38 mg, 0.05 mmol) in anhydrous toluene (8 mL) was subject to a freeze-pump-thaw cycle followed by backfilling with argon. The mixture was heated at reflux under argon in the dark for 21 h. When the reaction mixture had cooled to room temperature, diethyl ether (20 mL) was added and then mixture was washed with brine (50 mL). The isolated aqueous fraction was extracted with dichloromethane (3 × 30 mL). The combined organic fractions were dried over anhydrous sodium sulfate and filtered through a plug of Celite<sup>®</sup>. The filtrate was collected, and

the solvent removed. The crude product was purified in several steps: first, using size exclusive column chromatography over Biobeads<sup>TM</sup> with tetrahydrofuran as eluent; second, size exclusion column chromatography over Biobeads<sup>TM</sup> with toluene as eluent; third, using MPLC containing a C18 reverse-phase cartridge with methanol:acetonitrile mixtures as eluent (3:1 to 0:1); and finally, recrystallisation by dissolving the compound in diethyl ether followed by cooling the solution in a dry ice/2-propanol bath to afford **3** as a dark red colour powder (127 mg, 20%); m.p.: 75–76 °C;  $\lambda_{\max}$  (CH<sub>2</sub>Cl<sub>2</sub>/nm): 266 (log  $\epsilon$ /dm<sup>3</sup> mol<sup>-1</sup> cm<sup>-1</sup> 4.00), 311sh (3.75), 324 (3.85), 483 (4.60), 511sh (4.51);  $\delta_{\text{H}}$  (500 MHz, CDCl<sub>3</sub>): 2.29-2.34 (12H, m, Gl-CH<sub>2</sub>), 3.02-3.14 (12H, m, Gl-CH<sub>2</sub>), 3.30-3.41 (42H, m, Gl-CH<sub>2</sub> and -CH<sub>3</sub>), 6.97 (2H, d,  $J$  = 5.0 Hz, DTCPD-H), 7.05 (2H, s, DTCPD-H), 7.06 (2H, s, DTCPD-H), 7.18 (2H, d,  $J$  = 5.0 Hz, DTCPD-H);  $\delta_{\text{C}}$  (125 MHz, CDCl<sub>3</sub>): 37.5(8), 37.6(4), 49.6, 50.0, 59.0, 67.5(3), 67.5(5), 70.0, 70.1, 71.8, 117.3, 117.5, 121.7, 125.5, 134.9, 135.2, 136.4, 138.2, 138.7, 156.2, 156.8, 157.1;  $m/z$  (ESI-HRMS) anal. calcd. for C<sub>57</sub>H<sub>74</sub>O<sub>12</sub>S<sub>6</sub> [M]<sup>+</sup>: 1142.3499 (100%), 1143.3533 (62%), 1144.3457 (27%), 1145.3491 (17%), 1146.3524 (5%); found: 1142.3510 (56%), 1143.3564 (100%), 1144.3564 (70%), 1145.3557 (43%), 1146.3561 (19%).

**7,7'-[4,4,4',4',4'',4''-Hexakis(2-{2-methoxyethoxy}ethyl)-4*H*,4'*H*,4''*H*-(2,2':6',2''-tercyclopenta[2,1-*b*:3,4-*b'*]dithiophene) -6,6''-diyl]bis[benzo[*c*][1,2,5]thiadiazole-4-carbaldehyde] (4)**

A solution of **3** (100 mg, 0.09 mmol) in anhydrous tetrahydrofuran (10 mL) under argon was cooled in a dry ice/2-propanol bath. *n*-Butyllithium (2.5 M in hexanes, 0.1 mL) was added dropwise and then the reaction was stirred for 1 h with cooling in a dry ice/2-propanol bath. Tri-*n*-butyltin chloride (0.1 mL, 0.36 mmol) was added in one portion. The reaction solution was allowed to warm to room temperature and stirred for another 1.5 h. Deionised water (10 mL) was added to quench the reaction and the organic layer separated. The aqueous layer was extracted with diethyl ether (2 × 10 mL). The combined organic fractions were dried over anhydrous sodium sulfate, filtered, and the solvent was removed. The crude product was purified using size exclusion chromatography over BioBeads<sup>TM</sup> with tetrahydrofuran as eluent to afford the bis(tri-*n*-butylstannanyl) derivative as a dark red solid (70 mg), which was used immediately in the next reaction. A solution of the bis(tributylstannanyl) derivative (70 mg), 7-bromobenzo[*c*][1,2,5]thiadiazole-4-carbaldehyde (40 mg, 0.16 mmol) in anhydrous toluene (5 mL) was subject to a freeze-pump-thaw cycle followed by backfilling with argon. Tetrakis(triphenylphosphine)palladium(0) (5 mg, 0.004 mmol) was added, and the solution was heated at reflux under argon in an oil bath in the dark for 38 h. When the reaction mixture had cooled to room temperature, diethyl ether (20 mL) was added. The mixture was washed with brine (2 × 30 mL) and then the combined aqueous fractions were extracted with diethyl ether (30 mL). The combined organic fractions were dried over anhydrous sodium sulfate, filtered, and the solvent was removed. The crude product was purified by size exclusion column chromatography over Biobeads<sup>TM</sup> using a toluene:dichloromethane mixture (4:1) as eluent followed by reprecipitation from a mixture of dichloromethane:diethyl ether to afford **4** as a dark blue powder (51 mg, 40%); m.p.: 159–161 °C;  $\lambda_{\max}$  (CH<sub>2</sub>Cl<sub>2</sub>/nm): 321 (log  $\epsilon$ /dm<sup>3</sup> mol<sup>-1</sup> cm<sup>-1</sup> 4.14), 478 (4.32), 635 (4.72); IR (solid)  $\nu$ /cm<sup>-1</sup> = 1681 (C=O);  $\delta_{\text{H}}$  (500 MHz, CDCl<sub>3</sub>): 2.34-2.47 (12H, m, Gl-CH<sub>2</sub>), 3.14-3.20 (12H, m, Gl-CH<sub>2</sub>), 3.28 (12H, s, Gl-CH<sub>3</sub>), 3.32

(6H, s, GI-CH<sub>3</sub>), 3.35-3.42 (24H, m, GI-CH<sub>2</sub>), 7.14 (4H, brs, DTCPD-H), 7.98 (2H, brd, *J* = 8.0 Hz, BT-H), 8.22 (2H, d, *J* = 8.0 Hz, BT-H), 8.27 (2H, s, DTCPD-H), 10.69 (2H, s, CHO);  $\delta_{\text{C}}$  (125 MHz, CDCl<sub>3</sub>): 37.6, 50.2, 50.3, 58.9(9), 59.0(3), 67.5(2), 67.5(6), 70.1, 71.7(9), 71.8(2), 117.5, 118.1, 122.2, 124.6, 124.7, 133.0, 133.8, 134.8, 135.6, 138.6, 139.0, 141.0, 142.2, 152.0, 153.9, 157.4, 157.5, 159.4, 188.4; *m/z* (ESI-HRMS) anal. calcd. for C<sub>71</sub>H<sub>78</sub>N<sub>4</sub>O<sub>14</sub>S<sub>8</sub> [M]<sup>+</sup>: 1466.3275 (100%), 1467.3308 (77%), 1468.3233 (36%), 1469.3266 (28%); found: 1466.3285 (100%), 1467.3312 (86%), 1468.3296 (75%), 1469.3300 (43%), 1470.3286 (22%).

**2,2'-[({4,4',4'',4'''}-Hexakis[2-(2-methoxyethoxy)ethyl]-4*H*,4'*H*,4''*H*-[2,2':6',2''-tercyclopenta[2,1-*b*:3,4-*b'*]dithiophene]-6,6''-diyl}bis{benzo[*c*][1,2,5]thiadiazole-7,4-diyl}bis(methaneylylidene)]dimalononitrile (DTCPD-trimer)**

A mixture of **4** (40 mg, 0.03 mmol) and malononitrile (16 mg, 0.24 mmol) in anhydrous toluene (2 mL) was deoxygenated by placing under vacuum three times followed by backfilling with argon. Anhydrous pyridine (3 drops) was added to the reaction mixture before it was stirred in an oil bath held at 60 °C under argon for 5 h. When the reaction mixture was cooled to room temperature, dichloromethane (30 mL) was added, and the mixture was washed with brine (50 mL). The aqueous fraction was collected and extracted with dichloromethane (3 × 10 mL). The combined organic fractions were dried over anhydrous sodium sulfate, filtered, and the solvent was removed. The crude product was purified by size exclusive column chromatography over Biobeads<sup>TM</sup> using a toluene:dichloromethane mixture (3:1) as eluent followed by reprecipitation from a dichloromethane:diethyl ether mixture by cooling in a dry ice/acetone bath. The precipitate was collected at the filter, washed with cold diethyl ether (10 mL), and then dried under vacuum to afford **DTCPD-trimer** as a black powder (24 mg, 56%); m.p.: 190–191 °C;  $\lambda_{\text{max}}$  (CH<sub>2</sub>Cl<sub>2</sub>/nm): 259 (log  $\epsilon/\text{dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$  4.31), 334 (4.22), 399 sh (4.18), 482 (4.40), 726 (4.74);  $\lambda_{\text{max}}$  (film/nm): 341, 504, 799; IR (solid)  $\nu/\text{cm}^{-1}$  = 2221 (C≡N);  $\delta_{\text{H}}$  (500 MHz, CDCl<sub>3</sub>): 2.34-2.47 (12H, m, GI-CH<sub>2</sub>), 3.14-3.21 (12H, m, GI-CH<sub>2</sub>), 3.28 (12H, s, GI-CH<sub>3</sub>), 3.32 (6H, s, GI-CH<sub>3</sub>), 3.34-3.43 (24H, m, GI-CH<sub>2</sub>), 7.14 (2H, brs, DTCPD-H), 7.16 (2H, s, DTCPD-H), 7.97 (2H, brd, *J* = 8.0 Hz, BT-H), 8.32 (2H, s, DTCPD-H), 8.76 (2H, d, *J* = 8.0 Hz, BT-H), 8.77 (2H, s, VIN-H);  $\delta_{\text{C}}$  (125 MHz, CDCl<sub>3</sub>): 37.6, 50.2, 50.4, 59.0(1), 59.0(4), 67.5(2), 67.5(5), 70.2, 71.8(0), 71.8(3), 80.5, 113.6, 114.3, 117.6, 118.3, 120.5, 122.5, 125.6, 130.8, 134.1, 134.7, 135.9, 138.6, 139.2, 141.9, 143.8, 150.8, 151.9, 154.6, 157.6, 158.1, 160.3; *m/z* (ESI-HRMS) anal. calcd. for C<sub>77</sub>H<sub>78</sub>N<sub>8</sub>O<sub>12</sub>S<sub>8</sub> [M]<sup>+</sup>: 1562.3499 (100%), 1563.3533 (83%), 1564.3457 (36%), 1565.3491 (30%), 1566.3524 (12%); found: 1562.3532 (100%), 1563.3558 (92%), 1564.3548 (79%), 1565.3550 (46%), 1566.3540 (21%); *T*<sub>d(5%)</sub>: 316 °C; DSC (heating and cooling rate: 100 °C min<sup>-1</sup>): *T*<sub>m</sub> = 195 °C (1<sup>st</sup> cycle – heating), 175 °C (2<sup>nd</sup> and 3<sup>rd</sup> cycles – heating), not present in the 4<sup>th</sup> cycle heating; *T*<sub>g</sub> ≈ 93 °C (2<sup>nd</sup> and subsequent cycles – heating); CV: *E*<sub>1/2(ox)</sub> (dichloromethane): 0.05 V, 0.2 V, 0.9 V; *E*<sub>1/2(red)</sub> (tetrahydrofuran): -1.1 V, -1.7 V.

**7-[7-Bromo-9,9-bis(2-{2-methoxyethoxy}ethyl)-9*H*-fluoren-2-yl]benzo[*c*][1,2,5]thiadiazole-4-carbaldehyde (**5**)**

Tetrakis(triphenylphosphine)palladium(0) (117 mg, 0.10 mmol) followed by *t*-butanol (10 mL) were added to a mixture of 2-bromo-7-iodo-9,9-bis[2-(2-methoxyethoxy)ethyl]-9*H*-fluorene (see Supporting Information) (1.53 g, 2.7 mmol), 7-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)benzo[*c*][1,2,5]thiadiazole-4-carbaldehyde<sup>26</sup> (837 mg, 2.9 mmol), aqueous potassium carbonate (2 M, 20 mL), and toluene (25 mL) that had been deoxygenated by placing under vacuum six times and following by backfilling with argon. The mixture was stirred at reflux under argon overnight in the dark. The mixture was allowed to cool to room temperature and diluted with diethyl ether (20 mL) and water (20 mL). The aqueous layer was isolated and extracted with diethyl ether (5 × 20 mL). The combined organic fractions were washed with water (2 × 20 mL), brine (2 × 50 mL), dried over anhydrous sodium sulfate, and filtered through a silica plug that was pre-washed with diethyl ether (~100 mL). The filtrate was collected, and the solvent was removed. The crude product was purified using flash column chromatography over silica with diethyl ether:toluene mixtures (0:1 to 1:1) as eluent to afford **5** as a bright orange oil (843 mg, 52%); found: C, 58.9; H, 5.1; N, 5.0; S, 5.1; C<sub>30</sub>H<sub>31</sub>BrN<sub>2</sub>O<sub>5</sub>S requires: C, 58.9; H, 5.1; N, 4.6; S, 5.2; λ<sub>max</sub>(CH<sub>2</sub>Cl<sub>2</sub>/nm): 240sh (log ε/dm<sup>3</sup> mol<sup>-1</sup> cm<sup>-1</sup> 4.05), 301 (4.52), 310 sh (4.49), 405 (4.36); IR (solid) ν/cm<sup>-1</sup> = 1688 (C=O); δ<sub>H</sub> (500 MHz, CDCl<sub>3</sub>): 2.41–2.53 (4H, m, Gl–CH<sub>2</sub>), 2.84–2.93 (4H, m, Gl–CH<sub>2</sub>), 3.21–3.23 (4H, m, Gl–CH<sub>2</sub>), 3.26 (6H, s, Gl–CH<sub>3</sub>), 3.28–3.30 (4H, m, Gl–CH<sub>2</sub>), 7.53 (1H, dd, *J* = 2.0, 8.0 Hz, Fl–H), 7.62 (1H, d, *J* = 1.0 Hz Fl–H), 7.63 (1H, d, *J* = 8.5 Hz, Fl–H), 7.84 (1H, dd, *J* = 1.0, 8.0 Hz, Fl–H), 7.97 (1H, d, *J* = 7.5 Hz, BT–H), 8.05–8.08 (2H, m, Fl–H), 8.34 (1H, d, *J* = 7.5 Hz, BT–H), 10.82 (1H, s, CHO); δ<sub>C</sub> (125 MHz, CDCl<sub>3</sub>): 39.5, 51.9, 59.0, 67.0, 70.0, 71.7, 120.1, 121.6, 122.1, 124.6, 126.3, 126.9(0), 126.9(4), 129.4, 130.8, 132.6, 135.8, 138.7, 140.3, 140.9, 149.3, 151.8, 153.8, 154.0, 189.0; *m/z* (ESI-HRMS) calcd. for C<sub>30</sub>H<sub>31</sub>BrN<sub>2</sub>O<sub>5</sub>S: 611.1210 (92%), 612.1241 (32%), 613.1194 (100%), 614.1222 (33%); found: 611.1233 (96%), 612.1275 (29%), 613.1211 (100%), 614.1245 (33%).

**7,7'-[9,9,9',9'-Tetrakis(2-{2-methoxyethoxy}ethyl)-9*H*,9'*H*-(2,2'-bifluorene)-7,7'-diyl]bis[benzo[*c*][1,2,5]thiadiazole-4-carbaldehyde] (7)**

[1,1'-Bis(diphenylphosphino)ferrocene]dichloropalladium(II) dichloromethane complex (66 mg, 0.08 mmol) was added to a mixture of **5** (926 mg, 1.5 mmol), bis(pinacol)diboron (472 mg, 1.8 mmol), anhydrous potassium acetate (491 mg, 5.0 mmol), and dry 1,4-dioxane (14.5 mL) that had been deoxygenated by placing under vacuum three times followed by backfilling with argon. The mixture was stirred at reflux under argon in the dark overnight. The mixture was allowed to cool to room temperature and filtered through a Celite<sup>®</sup> plug that was pre-washed with diethyl ether. The residue was washed thoroughly with dichloromethane (~150 mL). The filtrate was collected, and the solvent was removed. The crude product was purified using flash column chromatography over silica with *n*-hexane:diethyl ether:dichloromethane mixtures (1:1:0 then 1:2:1) as eluent to afford **6** as a bright orange solid (663 mg), which was used directly in the following reaction without further purification. Tetrakis(triphenylphosphine)palladium(0) (10 mg, 0.01 mmol) was added to a mixture of **5** (171 mg, 0.28 mmol), **6** (205 mg), aqueous potassium carbonate (2 M, 0.4 mL), toluene (1.6 mL) and *t*-butanol (0.2 mL) that had been deoxygenated under vacuum three times followed by backfilling with

argon. The reaction mixture was stirred at reflux under argon in the dark for 40 h. The reaction was allowed to cool to room temperature before brine (20 mL) was added followed by dichloromethane (30 mL) and diethyl ether (60 mL). The aqueous layer was separated, and the organic fraction was washed with deionized water (3 × 30 mL). The organic fraction was then dried over anhydrous sodium sulfate, filtered, and the solvent was removed. The crude product was purified using MPLC over silica with a diethyl ether:methanol mixture (19:1) as eluent to afford an orange solid. The orange solid was dissolved in methanol (5 mL) and diethyl ether (45 mL) was added. The mixture was cooled in a dry ice/acetone bath and the precipitate was collected via vacuum filtration and dried under vacuum to afford **7** as a bright orange powder (170 mg, 59% based on the amount of **5** used in the second step); m.p.: 175–177 °C;  $\lambda_{\text{max}}$ (CH<sub>2</sub>Cl<sub>2</sub>/nm): 324 (log  $\epsilon/\text{dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$  5.11), 431 (5.01); IR (solid)  $\nu/\text{cm}^{-1}$  = 1685 (C=O);  $\delta_{\text{H}}$  (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 2.51-2.60 (8H, m, Gl-CH<sub>2</sub>), 2.92-3.00 (8H, m, Gl-CH<sub>2</sub>), 3.17 (12H, s, Gl-CH<sub>3</sub>), 3.20-3.23 (8H, m, Gl-CH<sub>2</sub>), 3.24-3.26 (8H, m, Gl-CH<sub>2</sub>), 7.79 (2H, dd,  $J$  = 1.5, 8.0 Hz, Fl-H), 7.85 (2H, brd,  $J$  = 1.0 Hz, Fl-H), 7.92 (2H, d,  $J$  = 8.0 Hz, Fl-H), 7.95 (2H, d,  $J$  = 8.0 Hz, Fl-H), 8.03 (2H, d,  $J$  = 7.5 Hz, BT-H), 8.12 (2H, dd,  $J$  = 1.5, 8.0 Hz, Fl-H), 8.19 (2H, brd,  $J$  = 1.0 Hz, Fl-H), 8.34 (2H, d,  $J$  = 7.5 Hz, BT-H), 10.82 (2H, s, CHO);  $\delta_{\text{C}}$  (125 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 39.7, 51.9, 58.6, 67.1, 69.9, 71.8, 120.2, 120.8, 121.9, 124.7, 126.3, 126.9, 127.1, 129.3, 132.2, 135.6, 139.3, 140.4, 141.1, 141.7, 150.1, 150.8, 154.1(3), 154.1(5), 188.9;  $m/z$  (ESI-HRMS) calcd. for C<sub>66</sub>H<sub>62</sub>N<sub>4</sub>O<sub>10</sub>S<sub>2</sub>Na [M + Na]<sup>+</sup>: 1085.3800 (100%), 1086.3833 (65%), 1087.3867 (21%); found: 1085.3767 (100%), 1086.3799 (68%), 1087.3810 (31%).

**2,2'-[(9,9,9',9'-Tetrakis[2-(2-methoxyethoxy)ethyl]-9H,9'H-[2,2'-bifluorene]-7,7'-diyl]bis[benzo[*c*][1,2,5]thiadiazole-7,4-diyl]bis(methanelylidene)]dimalononitrile (Fluorenyl-dimer)**

A mixture of **7** (53 mg, 0.05 mmol) and malononitrile (35 mg, 0.53 mmol) in chlorobenzene (2 mL) was deoxygenated by placing under vacuum three times followed by backfilling with argon. The reaction mixture was then stirred in an oil bath held at 75 °C under argon. Pyridine (0.1 mL) was added to the reaction mixture, which was then stirred in an oil bath held at 75 °C under argon overnight. The solvent was removed and the crude product was purified using MPLC over silica with a diethyl ether:methanol mixture (19:1) as eluent to afford a dark red solid. The solid was dissolved in ethyl acetate (5 mL) and then *n*-hexane (30 mL) was added. The mixture was cooled in a dry ice/acetone bath and the precipitate that formed was collected at the filter, which was then washed with *n*-hexane (20 mL), and dried under vacuum to afford **Fluorenyl-dimer** as a bright red solid (20 mg, 37%); m.p.: 71 °C (DSC);  $\lambda_{\text{max}}$ (CH<sub>2</sub>Cl<sub>2</sub>/nm): 275 (log  $\epsilon/\text{dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$  5.16), 337 (4.70), 480 (4.66); IR (solid)  $\nu/\text{cm}^{-1}$  = 2226 (C≡N);  $\delta_{\text{H}}$  (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 2.51-2.60 (8H, m, Gl-CH<sub>2</sub>), 2.91-3.00 (8H, m, Gl-CH<sub>2</sub>), 3.17 (12H, s, Gl-CH<sub>3</sub>), 3.20-3.22 (8H, m, Gl-CH<sub>2</sub>), 3.24-3.26 (8H, m, Gl-CH<sub>2</sub>), 7.79 (2H, dd,  $J$  = 1.5, 8.0 Hz, Fl-H), 7.85 (2H, d,  $J$  = 1.5 Hz, Fl-H), 7.92 (2H, d,  $J$  = 8.0 Hz, Fl-H), 7.96 (2H, d,  $J$  = 8.0 Hz, Fl-H), 8.06 (2H, d,  $J$  = 7.5 Hz, BT-H), 8.14 (2H, dd,  $J$  = 1.5, 8.0 Hz, Fl-H), 8.24 (2H, d,  $J$  = 1.5 Hz, Fl-H), 8.84 (2H, d,  $J$  = 7.5 Hz, BT-H), 8.91 (2H, s, VIN-H);  $\delta_{\text{C}}$  (125 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 39.7, 51.9, 58.6, 67.1, 69.9, 71.8, 83.3, 113.2, 114.0, 120.3, 120.9, 121.9, 122.6, 124.8, 126.9, 127.4, 129.4, 130.8, 135.2, 139.2, 140.5, 141.2, 142.1, 150.2, 150.9, 153.0, 153.2, 154.6;  $m/z$  (ESI-HRMS) calcd. for

$C_{66}H_{63}N_8O_8S_2$  [M + H]<sup>+</sup>: 1159.4205 (100%), 1160.4238 (71%), 1161.4272 (25%); found: 1159.4224 (100%), 1160.4252 (73%), 1161.4266 (38%);  $T_{d(5\%)}$ : 336 °C; DSC (heating rate: 50 °C min<sup>-1</sup>/cooling rate: 100 °C):  $T_m$  = 71 °C (1<sup>st</sup> cycle – heating),  $T_g$  = 66 °C (2<sup>nd</sup> – 4<sup>th</sup> cycles – heating);  $E_{1/2(\text{red})}$  (CV/tetrahydrofuran): -1.2 V, -1.8 V.

**7,7'-[9,9,9',9',9'',9''-Hexakis(2-{2-methoxyethoxy}ethyl)-9H,9'H,9''H-(2,2':7',2''-terfluorene)-7,7''-diyl]bis[benzo[c][1,2,5]thiadiazole-4-carbaldehyde] (9)**

Tetrakis(triphenylphosphine)palladium(0) (15 mg, 0.01 mmol) was added to a mixture of **8**<sup>[29]</sup> (52 mg, 0.10 mmol), crude **6** (151 mg), caesium carbonate (99 mg, 0.30 mmol), and anhydrous toluene (1.5 mL) that had been deoxygenated by three freeze-pump-thaw cycles followed by backfilling with argon. The reaction mixture was stirred at reflux under argon in the dark for 22 h. The reaction was allowed to cool to room temperature before deionised water (10 mL) and dichloromethane (10 mL) were added sequentially. The aqueous layer was separated and extracted with dichloromethane (3 × 10 mL). The combined organic fractions were dried over anhydrous sodium sulfate, filtered, and the solvent was removed. The crude product was purified using MPLC over silica using diethyl ether:methanol mixtures (100:0 then increasing to 93:7) as eluent to afford an orange solid. The solid was dissolved in diethyl ether (15 mL) and then the solution was cooled in a dry ice/acetone bath. The precipitate was collected via vacuum filtration and dried under vacuum to afford **9** as a bright orange powder (50 mg, 35%); m.p.: 68–69 °C;  $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2/\text{nm})$ : 330 (log  $\epsilon/\text{dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$  4.78), 349 (4.81), 425 (4.74); IR (solid)  $\nu/\text{cm}^{-1}$  = 1689 (C=O);  $\delta_{\text{H}}$  (300 MHz,  $\text{CDCl}_3$ ): 2.58-2.66 (12H, m, Gl-CH<sub>2</sub>), 2.90-3.01 (12H, m, Gl-CH<sub>2</sub>), 3.24-3.35 (42H, m, Gl-CH<sub>2</sub> and -CH<sub>3</sub>), 7.70-7.78 (8H, m, Fl-H), 7.84 (2H, d,  $J$  = 8.0 Hz, Fl-H), 7.88 (2H, d,  $J$  = 8.0 Hz, Fl-H), 7.92-7.95 (2H, m, Fl-H), 8.01 (2H, d,  $J$  = 7.5 Hz, BT-H), 8.10-8.13 (4H, m, Fl-H), 8.36 (2H, d,  $J$  = 7.5 Hz, BT-H), 10.83 (2H, s, CHO);  $m/z$  (ESI-HRMS) calcd. for  $C_{83}H_{90}N_4O_{14}S_2$  [M]<sup>+</sup>: 1430.5889 (100%), 1431.5923 (90%), 1432.5957 (40%), 1433.5990 (12%); found: 1430.5894 (37%), 1431.5957 (100%), 1432.5994 (80%), 1433.6012 (46%).

**2,2'-[({9,9,9',9',9'',9''-Hexakis[2-(2-methoxyethoxy)ethyl]-9H,9'H,9''H-[2,2':7',2''-terfluorene]-7,7''-diyl})bis{benzo[c][1,2,5]thiadiazole-7,4-diyl})bis(methaneylylidene)]dimalononitrile (Fluorenyl-trimer)**

A mixture of **9** (50 mg, 0.03 mmol) and malononitrile (35 mg, 0.53 mmol) in anhydrous chloroform (1.5 mL) was deoxygenated by placing under vacuum three times followed by backfilling with argon. Pyridine (0.1 mL) was added to the reaction mixture before it was stirred in an oil bath held at 75 °C under argon for 3.5 h. When the reaction mixture had cooled to room temperature, dichloromethane (30 mL) was added, and the mixture was washed with deionised water (30 mL). The aqueous fraction was then extracted with dichloromethane (3 × 10 mL). The combined organic fractions were dried over anhydrous sodium sulfate, filtered, and the solvent was removed under a stream of nitrogen. The crude product was purified by recrystallisation by dissolving in methanol followed by cooling the solution in a dry ice/acetone bath. The precipitate formed was collected at the filter, washed with cold methanol (10 mL), and dried under vacuum to afford **Fluorenyl-trimer** as a dark red powder (33 mg, 62%);  $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2/\text{nm})$ : 350 (log  $\epsilon/\text{dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$

<sup>1</sup> 4.78), 479 (4.56); IR (solid)  $\nu/\text{cm}^{-1} = 2226$  (C≡N);  $\delta_{\text{H}}$  (500 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 2.53-2.59 (12H, m, GI-CH<sub>2</sub>), 2.89-3.01 (12H, m, GI-CH<sub>2</sub>), 3.18-3.27 (42H, m, GI-CH<sub>2</sub> and -CH<sub>3</sub>), 7.77 (2H, dd,  $J = 1.5, 8.0$  Hz, FI-H), 7.79 (2H, dd,  $J = 1.5, 8.0$  Hz, FI-H), 7.84 (4H, brdd,  $J = 1.0, 6.5$  Hz, FI-H), 7.87 (2H, d,  $J = 8.0$  Hz, FI-H), 7.92 (2H, d,  $J = 8.0$  Hz, FI-H), 7.94-7.96 (2H, m, FI-H), 8.06 (2H, d,  $J = 7.5$  Hz, BT-H), 8.14 (2H, dd,  $J = 1.5, 8.0$  Hz, FI-H), 8.24 (2H, brd,  $J = 1.0$  Hz, FI-H), 8.84 (2H, dd,  $J = 0.5, 7.5$  Hz, BT-H), 8.91 (2H, s, VIN-H);  $\delta_{\text{C}}$  (125 MHz, CD<sub>2</sub>Cl<sub>2</sub>): 40.0, 40.2, 52.0, 52.2, 58.9, 67.4, 70.2, 72.1, 72.2, 83.6, 113.5, 120.6, 120.7, 121.2, 122.1, 122.2, 122.9, 125.1, 127.1, 127.2, 127.7, 129.7, 131.1, 135.4, 139.3, 140.0, 140.7, 140.8, 141.7, 142.5, 150.6, 150.7, 151.2, 153.3, 153.5, 154.9;  $m/z$  (ESI-HRMS) calcd. for C<sub>89</sub>H<sub>90</sub>N<sub>8</sub>O<sub>12</sub>S<sub>2</sub>Na [M + Na]<sup>+</sup>: 1549.6012 (100%), 1550.6045 (96%), 1551.6079 (46%), 1552.6112 (14%); found: 1549.5993 (100%), 1550.6027 (99%), 1551.6053 (62%), 1552.6078 (31%);  $T_{\text{d}(5\%)}$ : 345 °C; DSC (heating/cooling rate: 100 °C min<sup>-1</sup>):  $T_{\text{g}} = 77$  °C (heating – third cycle), 191 °C (reversible endothermic transition – first and subsequent cycles);  $E_{1/2(\text{red})}$  (CV/tetrahydrofuran): -1.2 V, -1.8 V.

## Conflicts of interest

There are no conflicts to declare.

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## References

- [1] Y. Cui, H. Yao, J. Zhang, K. Xian, T. Zhang, L. Hong, Y. Wang, Y. Xu, K. Ma, C. An, C. He, Z. Wei, F. Gao, J. Hou, *Adv. Mater.* **2020**, *32*, 1908205.

- [2] L. Hong, H. Yao, Z. Wu, Y. Cui, T. Zhang, Y. Xu, R. Yu, Q. Liao, B. Gao, K. Xian, H. Y. Woo, Z. Ge, J. Hou, *Adv. Mater.* **2019**, *31*, 1903441.
- [3] B. Kan, J. Zhang, F. Liu, X. Wan, C. Li, X. Ke, Y. Wang, H. Feng, Y. Zhang, G. Long, R. H. Friend, A. A. Bakulin, Y. Chen, *Adv. Mater.* **2018**, *30*, 1704904.
- [4] C. Zhu, J. Yuan, F. Cai, L. Meng, H. Zhang, H. Chen, J. Li, B. Qiu, H. Peng, S. Chen, Y. Hu, C. Yang, F. Gao, Y. Zou, Y. Li, *Energy & Environ. Sci.* **2020**, *13*, 2459.
- [5] L. Zuo, X. Shi, S. B. Jo, Y. Liu, F. Lin, A. K.-Y. Jen, *Adv. Mater.* **2018**, *30*, 1706816.
- [6] L. Zhu, M. Zhang, G. Zhou, T. Hao, J. Xu, J. Wang, C. Qiu, N. Prine, J. Ali, W. Feng, X. Gu, Z. Ma, Z. Tang, H. Zhu, L. Ying, Y. Zhang, F. Liu, *Adv. Energy Mater.* **2020**, *10*, 1904234.
- [7] X. Song, N. Gasparini, L. Ye, H. Yao, J. Hou, H. Ade, D. Baran, *ACS Energy Lett.* **2018**, *3*, 669.
- [8] R. Xue, J. Zhang, Y. Li, Y. Li, *Small* **2018**, *14*, 1801793.
- [9] S. Lucas, J. Kammerer, M. Pfannmöller, R. R. Schröder, Y. He, N. Li, C. J. Brabec, T. Leydecker, P. Samorì, T. Marszalek, W. Pisula, E. Mena-Osteritz, P. Bäuerle, *Sol. RRL* **2021**, *5*, 2000653.
- [10] W. Wang, R. Sun, J. Guo, J. Guo, J. Min, *Angew. Chem. Int. Ed.* **2019**, *58*, 14556.
- [11] Y. He, T. Heumüller, W. Lai, G. Feng, A. Classen, X. Du, C. Liu, W. Li, N. Li, C. J. Brabec, *Adv. Energy Mater.* **2019**, *9*, 1900409.
- [12] G. Feng, J. Li, Y. He, W. Zheng, J. Wang, C. Li, Z. Tang, A. Osvet, N. Li, C. J. Brabec, Y. Yi, H. Yan, W. Li, *Joule* **2019**, *3*, 1765.
- [13] Q. Lin, A. Armin, R. C. R. Nagiri, P. L. Burn, P. Meredith, *Nat. Photon.* **2015**, *9*, 106.
- [14] S. Collavini, S. F. Völker, J. L. Delgado, *Angew. Chem. Int. Ed.* **2015**, *54*, 9757.
- [15] J. E. Donaghey, A. Armin, D. M. Stoltzfus, P. L. Burn, P. Meredith, *Chem. Comm.* **2016**, *52*, 13714.
- [16] M. P. Hughes, K. D. Rosenthal, N. A. Ran, M. Seifrid, G. C. Bazan, T.-Q. Nguyen, *Adv. Funct. Mater.* **2018**, *28*, 1801542.
- [17] X. Chen, Z. Zhang, Z. Ding, J. Liu, L. Wang, *Angew. Chem. Int. Ed.* **2016**, *55*, 10376.
- [18] S. Torabi, F. Jahani, I. Van Severen, C. Kanimozhi, S. Patil, R. W. A. Havenith, R. C. Chiechi, L. Lutsen, D. J. M. Vanderzande, T. J. Cleij, J. C. Hummelen, L. J. A. Koster, *Adv. Funct. Mater.* **2015**, *25*, 150.
- [19] A. Armin, D. M. Stoltzfus, J. E. Donaghey, A. J. Clulow, R. C. R. Nagiri, P. L. Burn, I. R. Gentle, P. Meredith, *J. Mater. Chem. C* **2017**, *5*, 3736.
- [20] S. Sami, R. Alessandri, R. Broer, R. W. A. Havenith, *ACS Appl. Mater. Interfaces* **2020**, *12*, 17783.
- [21] S. Kraner, C. Koerner, K. Leo, E. Bittrich, K. J. Eichhorn, Y. Karpov, A. Kiriy, M. Stamm, K. Hinrichs, M. Al-Hussein, *Phys. Rev. B* **2015**, *91*, 195202.
- [22] B. Meng, H. Song, X. Chen, Z. Xie, J. Liu, L. Wang, *Macromolecules* **2015**, *48*, 4357.
- [23] A. Wadsworth, C. Nielsen, S. Holliday, I. McCulloch, M. Kirkus, D. Baran, S. Ashraf, International Patent WO2017/191644 A1, May 6, **2016**.
- [24] C. W. Tang, *Appl. Phys. Lett.* **1986**, *48*, 183.



- [25] J. Li, F. Dierschke, J. Wu, A. C. Grimsdale, K. Müllen, *J. Mater. Chem.* **2006**, *16*, 96.
- [26] R. Y. C. Shin, T. Kietzke, S. Sudhakar, A. Dodabalapur, Z.-K. Chen, A. Sellinger, *Chem. Mater.* **2007**, *19*, 1892.
- [27] N. M. Kirby, S. T. Mudie, A. M. Hawley, D. J. Cookson, H. D. T. Mertens, N. Cowieson, V. Samardzic-Boban, *J. Appl. Cryst.* **2013**, *46*, 1670.
- [28] J. Ilavsky, *J. Appl. Cryst.* **2012**, *45*, 324.
- [29] S. Gug, F. Bolze, A. Specht, S. Bourgogne, M. Goeldner, J-F. Nicoud, *Angew. Chem. Int. Ed.* **2008**, *47*, 9525.