



Cite this: *J. Mater. Chem. C*, 2017, 5, 1997

## Thieno[3,4-c]pyrrole-4,6-dione as novel building block for host materials for red PhOLEDs†

Paul Kautny,<sup>a</sup> Chenyang Zhao,<sup>b</sup> Dominik Schopf,<sup>a</sup> Berthold Stöger,<sup>c</sup> Ernst Horkel,<sup>a</sup> Jiangshan Chen,<sup>\*b</sup> Dongge Ma,<sup>b</sup> Johannes Fröhlich<sup>a</sup> and Daniel Lumpi<sup>a</sup>

In the presented work the electron accepting thieno[3,4-c]pyrrole-4,6-dione is introduced as a novel building block for donor–acceptor based host materials for Phosphorescent Organic Light Emitting Diodes (PhOLEDs). A series of three regioisomers consisting of the thieno[3,4-c]pyrrole-4,6-dione acceptor and carbazole donors linked via a phenylene linker was prepared and the impact of the phenylene substitution pattern on the molecular properties was analyzed. Regarding their applicability as host materials, the newly developed materials were investigated in red PhOLED devices achieving a high current efficiency of 30.6 cd A<sup>−1</sup> corresponding to an external quantum efficiency of 17.7% and a high power efficiency of 23.8 lm W<sup>−1</sup>. Thus, we present the first successful application of the thieno[3,4-c]pyrrole-4,6-dione building block in host materials for PhOLEDs.

Received 3rd December 2016,  
Accepted 31st January 2017

DOI: 10.1039/c6tc05248g

rsc.li/materials-c

### Introduction

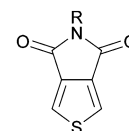
Since the first reports on electroluminescence from organic materials,<sup>1,2</sup> great efforts have been made in the development of organic light emitting diodes (OLEDs).<sup>3–8</sup> The application of transition metal emitters in phosphorescent OLEDs (PhOLEDs)<sup>9,10</sup> and of organic compounds capable of thermally activated delayed fluorescence (TADF)<sup>11</sup> boosted the efficiency of OLED devices.<sup>12–16</sup> Therefore, one major goal of research is the development of new host materials for these emitters.<sup>17–21</sup> In particular, the application of bipolar host materials consisting of donor and acceptor subunits proved successful, owing to their balanced charge transport properties and thus simplified device structures and broad recombination zones.<sup>22–24</sup>

The primary source of electron donating groups for optoelectronic materials are aromatic amines, such as triphenylamines,<sup>3,9,25–29</sup> carbazoles,<sup>10,17,28,30–33</sup> indolocarbazoles,<sup>34–37</sup> phenoxazines<sup>15,38–40</sup> and dihydroacridines.<sup>15,41</sup> In contrast, the nature of the electron accepting moieties exhibits more variations. Among the most frequently used acceptors are oxadiazoles,<sup>27,33,38,42–44</sup> phenantrolines,<sup>31,45</sup> pyridines,<sup>32,46</sup>

benzimidazoles,<sup>26,28,47</sup> 1,3,5-triazines,<sup>39,48,49</sup> 1,2,4-triazoles,<sup>29,38,50</sup> phosphine oxides,<sup>51–53</sup> sulfones,<sup>15,54</sup> ketones,<sup>40,41</sup> carbolines<sup>55,56</sup> and benzonitriles.<sup>11,57</sup>

The thieno[3,4-c]pyrrole-4,6-dione (TPD) (Scheme 1) has been successfully employed as electron accepting unit in donor–acceptor polymers for organic photovoltaics in combination with various donor units.<sup>58–62</sup> Moreover, the application of a TPD based polymer in an organic field effect transistor has been reported.<sup>63</sup> So far however, this particular unit has not been exploited as acceptor in materials for OLEDs.

This circumstance prompted us to explore the possibility to utilize the TPD building block in donor–acceptor host materials for PhOLEDs. Hence, this study presents the first application of TPD in host materials in combination with the well-established carbazole donor. The variation of the linkage mode between the two molecular subunits allowed for a modification of the molecular properties of the derivatives. Suchlike developed compounds were employed as host materials in efficient red PhOLED devices (CE<sub>max</sub>: 30.6 cd A<sup>−1</sup>, PE<sub>max</sub>: 23.8 lm W<sup>−1</sup>, EQE<sub>max</sub>: 17.7%), demonstrating the applicability of the TPD



Thieno[3,4-c]pyrrole-4,6-dione

TPD

Scheme 1 Molecular structure of thieno[3,4-c]pyrrole-4,6-dione (TPD).

<sup>a</sup> Institute of Applied Synthetic Chemistry, TU Wien, Getreidemarkt 9/163, A-1060 Vienna, Austria. E-mail: paul.kautny@tuwien.ac.at

<sup>b</sup> State Key Laboratory of Polymer Physics and Chemistry, Changchun Institute of Applied Chemistry, Chinese Academy of Sciences, Changchun, 130022, China. E-mail: jschen@ciac.ac.cn

<sup>c</sup> Institute of Chemical Technologies and Analytics, TU Wien, Getreidemarkt 9/164, A-1060 Vienna, Austria

† Electronic supplementary information (ESI) available: NMR spectra, DSC, TGA and CV analysis and crystallographic information. CCDC 1519402. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6tc05248g

unit as electron withdrawing organic building block for optoelectronic applications.

## Results and discussion

### Synthesis

Three different TPD based host materials (*p*-PCzTPD, *m*-PCzTPD and *o*-PCzTPD), in which the TPD unit is connected to the carbazole donors by phenylene linkers, were prepared (Scheme 2). Thereby three different linkage modes (*para*, *meta*, *ortho*) were realized using the phenylene linker. The unsubstituted imide of the TPD unit was protected with an octyl chain, also facilitating the handling of the materials during the synthetic process. The key synthetic step in the preparation of the new materials was a Suzuki cross coupling reaction of dibrominated TPD **1** and the suitable substituted boronic acid esters **2a–c** (Scheme 2). Products *p*-PCzTPD and *m*-PCzTPD were obtained with yields of 67% and 70%, respectively. In contrast the reaction towards *o*-PCzTPD proceeded significantly slower and with a low yield of 17% due to the high steric demand resulting from the *ortho* linkage mode. All materials were characterized by  $^1\text{H}$  and  $^{13}\text{C}$  NMR and high resolution mass spectrometry.

In the case of *p*-PCzTPD the molecular structure was unequivocally confirmed by X-ray crystal analysis (Fig. 1). Both phenylene linkers are close to coplanar with the TPD core [angles between the least square (LS) planes:  $\Phi_2 = 15.04(11)^\circ$  and  $\Phi_3 = 18.08(11)^\circ$ ] and thus fully conjugated with the central aromatic system. In contrast, the carbazole moieties and the phenylene linkers are more pronouncedly inclined [angles between the LS planes:  $\Phi_1 = 54.53(12)^\circ$  and  $\Phi_4 = 61.68(11)^\circ$ ] due to the steric interaction of the *ortho* hydrogen atoms.

### Thermal properties

The thermal behaviour of the developed materials was studied to evaluate their applicability as host materials for PhOLEDs (Table 1). *m*-PCzTPD and *p*-PCzTPD exhibited high decomposition temperatures ( $T_d$  – corresponding to 5% mass loss) of 438 °C and 448 °C during thermogravimetric analysis (Fig. S7, ESI†). In contrast, the  $T_d$  of *o*-PCzTPD was lower (369 °C). The glass transition temperatures ( $T_g$ ) of the compounds were determined by differential scanning calorimetry, to estimate the morphological stability of thin films of the materials. Whereas *o*-PCzTPD exhibited a rather low  $T_g$  of 62 °C, the  $T_g$ s increased with increasingly linear

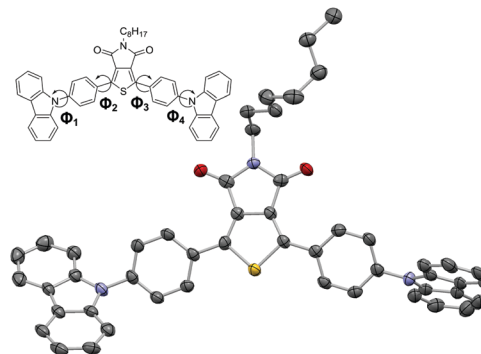


Fig. 1 Molecular structure of *p*-PCzTPD. C, N, O and S atoms are represented by gray, blue, red and yellow ellipsoids drawn at 70% probability levels. H atoms are omitted for clarity.

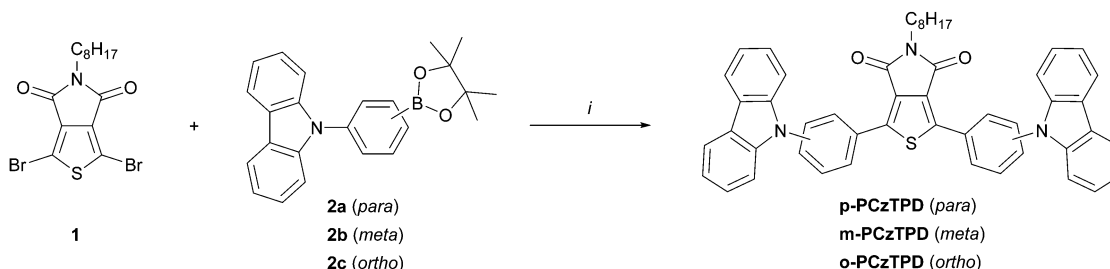
shape of the molecules to 84 °C for *m*-PCzTPD and 102 °C for *p*-PCzTPD (Fig. S8, ESI†).

### Photophysical properties

UV/Vis absorption, room temperature photoluminescence and low temperature phosphorescence spectra were recorded to investigate the impact of the substitution pattern on the photophysical properties of the materials (Fig. 2). The key photophysical data is summarized in Table 1. All derivatives exhibit a distinct absorption peak at 292 nm, which can be attributed to a transition centered on the carbazole moieties.<sup>31</sup> Moreover, weaker transitions at approximately 325 nm and 340 nm, which are typical for phenylcarbazole derivatives,<sup>64</sup> are found for all compounds. Fully conjugated *p*-PCzTPD features an additional broad absorption peak with a peak maximum at 386 nm. This peak can be attributed to an intramolecular charge transfer transition between the electron rich carbazoles and the TPD acceptor unit. Owing to the decreased effective conjugation resulting from the *ortho* and *meta* linkages,<sup>33</sup> this transition is absent in *o*-PCzTPD and *m*-PCzTPD. In analogy, the absorption onsets are shifted towards lower energy from *o*-PCzTPD (3.34 eV) to *m*-PCzTPD (3.18 eV) and *p*-PCzTPD (2.84 eV).

The fluorescence of the materials is distinctly red shifted compared to the absorption, and emission maxima are located at 476 nm (*p*-PCzTPD), 506 nm (*m*-PCzTPD) and 485 nm (*o*-PCzTPD).

To determine the triplet energy ( $E_T$ ) of the developed materials low temperature phosphorescence was recorded. The order of the



Scheme 2 Synthesis of TPD based host materials; (i) **1** (1.0 eq., 25 mM), **2a–c** (2.5 eq.), Pd(PPh<sub>3</sub>)<sub>4</sub> (0.05 eq.), K<sub>2</sub>CO<sub>3</sub> (5.0 eq., 2 M aqueous solution), THF, reflux.



Table 1 Physical data of the synthesized host materials

	$T_g/T_m/T_d^a$ [°C]	Opt. BG <sup>b,c</sup> [eV]	$\lambda_{PL,max}^c$ [nm]	HOMO/LUMO [eV]		$E_T$ [eV]	
				Exp. <sup>d</sup>	Cal. <sup>e</sup>	Exp. <sup>f</sup>	Cal. <sup>g</sup>
<b><i>p</i>-PCzTPD</b>	102/208/448	2.84	476	−5.68/−2.84	−5.68/−2.57	2.09	2.11
<b><i>m</i>-PCzTPD</b>	84/195/438	3.18	506	−5.70/−2.52	−5.71/−2.56	2.21	2.31
<b><i>o</i>-PCzTPD</b>	62/133/369	3.34	485	−5.65/−2.31	−5.75/−2.13	2.33	2.62

<sup>a</sup> Determined by thermogravimetric analysis and differential scanning calorimetry.  $T_m$ : melting temperature. <sup>b</sup> Estimated from the absorption onset. <sup>c</sup> Measurements were carried out in  $CH_2Cl_2$  (5  $\mu$ M) at room temperature. <sup>d</sup> HOMO levels were determined from the onset of the oxidation peak obtained during cyclic voltammetric measurements (0.5 mM solution in anhydrous  $CH_2Cl_2$  with  $Bu_4NBF_4$  (0.1 M) as supporting electrolyte); LUMO levels were calculated from the HOMO levels and the optical bandgap. <sup>e</sup> Calculated applying density functional theory level (B3LYP/6-311+G\*). <sup>f</sup> Estimated from the highest energy vibronic transition in solid solution of toluene/EtOH (9:1) at 77 K. <sup>g</sup> Calculated applying time dependent density functional theory level (B3LYP/6-311+G\*).

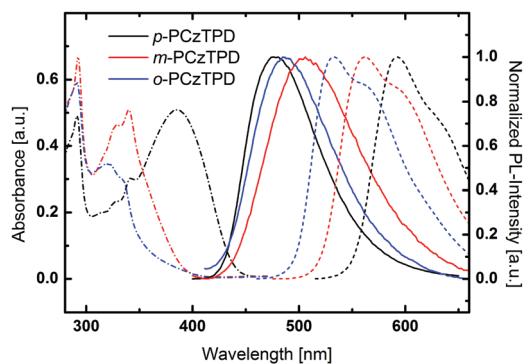


Fig. 2 UV-Vis absorption (dashed and dotted lines), normalized fluorescence at room temperature (solid lines) and normalized phosphorescence spectra at 77 K (dashed lines) of the materials under investigation.

emission maxima follows the absorption onset. Thus, ***o*-PCzTPD** features the highest  $E_T$  of 2.33 eV, whereas the  $E_T$  of ***m*-PCzTPD** is slightly lower (2.21 eV) and ***p*-PCzTPD** exhibits the lowest  $E_T$  of 2.09 eV. In ***p*-PCzTPD**, the pronounced donor-acceptor interaction between the carbazoles and the TPD, which is also evident in the absorption behaviour of the material, lowers the  $E_T$ s via intramolecular charge transfer.<sup>22</sup> The theoretically calculated  $E_T$  display the same trend, the absolute values for ***m*-PCzTPD** and ***o*-PCzTPD** are, however, higher than the experimentally determined energies. A closer inspection of the  $S_0 \rightarrow T_1$  transition reveals that in case of ***m*-PCzTPD** and ***o*-PCzTPD** transitions from low lying molecular orbitals contribute significantly to the lowest excited triplet state. These transitions may be slightly overestimated in the applied model, leading to the prediction of too high  $E_T$ s of the two derivatives. Ultimately, the application of different linkage modes allowed to control the degree of the overall conjugation in ***m*-PCzTPD** and ***o*-PCzTPD**. Therefore, the intramolecular charge transfer could be decreased, resulting in higher  $E_T$ s. Notably, in this series the *ortho* derivative ***o*-PCzTPD** exhibits a higher  $E_T$  than *meta* linked ***m*-PCzTPD**, which is unlike the behaviour of related host materials based on 1,3,4-oxadiazole acceptor units.<sup>33,65</sup>

### Electrochemical properties

The levels of the Highest Occupied Molecular Orbitals (HOMOs) of the materials were determined by cyclic voltammetry (CV). All derivatives exhibited irreversible oxidation, as commonly observed for carbazole based materials, owing to the instability of the

formed cations.<sup>66</sup> The HOMO levels were calculated from the onset of the oxidation peaks relative to the oxidation of ferrocene and span a narrow range between −5.65 eV and −5.70 eV (Fig. S9, ESI†), indicating no significant barrier for hole injection from the adjacent layers. The Lowest Unoccupied Molecular Orbital (LUMO) levels of the materials were estimated from the HOMO levels and the optical bandgaps and are located between −2.84 eV and −2.31 eV, whereby ***p*-PCzTPD** features the lowest value owing to the lower bandgap of this compound.

### Computational investigations

Density functional theory was applied to obtain a deeper understanding of the electronic layouts of the investigated materials on the molecular level. In general, the calculated HOMO and LUMO energy values are in good agreement with the experimental results (Table 1). Moreover, the energetic order of the triplet energies of the developed host materials was reliably reproduced by the theoretical calculations, albeit slightly higher energies were predicted for ***m*-PCzTPD** and ***o*-PCzTPD** compared to the experimental values.

The spatial distributions of the HOMOs and LUMOs are depicted in Fig. 3. In the case of the fully conjugated ***p*-PCzTPD**, the HOMO level expands over the whole aromatic backbone of the molecule with a distinctive localization of electron density on the central TPD unit. The LUMO of ***p*-PCzTPD** is more localized with major contributions coming from the TPD acceptor and the

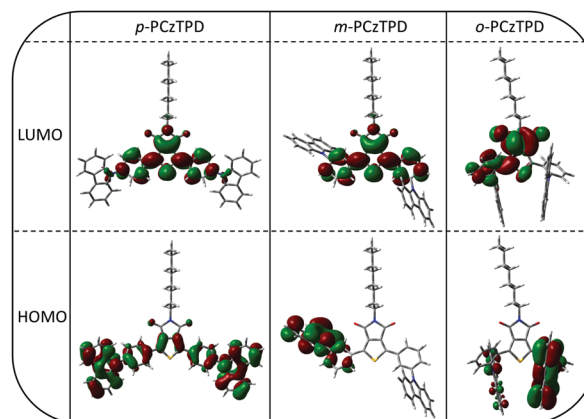


Fig. 3 Spatial distribution of the HOMOs and LUMOs of ***p*-PCzTPD**, ***m*-PCzTPD** and ***o*-PCzTPD**.



adjacent phenyl moieties. Looking at *m*-PCzTPD, the morphology of the HOMO is distinctly different. The electron density of the orbital is solely located on one of the carbazole units and the neighbouring phenyl ring owing to the decreased overall conjugation. In contrast, the shape of the LUMO is essentially unaffected by the altered linkage mode of the donor units. Consequently, the decreased spatial overlap of the two molecular orbitals leads to a lower degree of intramolecular charge transfer in *m*-PCzTPD, as reflected by the absence of a distinct charge transfer transition in the absorption spectra and also an increased  $E_T$ .<sup>22</sup>

In the case of *o*-PCzTPD, however, the overall situation changed as a result of the geometric structure of the molecule. The calculated shape of the aromatic part of *p*-PCzTPD is in excellent agreement with the molecular structure determined by X-ray diffraction. Accordingly, the optimized torsion angles between the aromatic subunits deviate only slightly from the experimentally determined values (Table 2). Similar torsion angles were observed for *m*-PCzTPD. In contrast the molecular structure of *o*-PCzTPD is distinctly different. Both carbazole units are oriented away from the TPD, owing to the steric demand of the carbonyl oxygen atoms (Fig. 3). This leads to a rotation of the phenylene linkers to avoid a close proximity of the carbazoles. Therefore, the torsion angles  $\Phi_2$  and  $\Phi_3$  between the TPD and the adjacent phenyl rings are distinctly enlarged to 49.43° and 53.46° and also the torsion angles between the phenylene linkers and the carbazole are larger than in the other two derivatives (Table 2). The highly twisted structure of *o*-PCzTPD restricts the HOMO to one of the carbazoles with only minor electron density on the neighbouring phenylene linker

as well as on the opposite carbazole. Notably, also the distribution of the LUMO is altered compared to the other derivatives. The LUMO is exclusively located on the TPD acceptor and the phenylene linker opposite of the carbazole that is hosting the HOMO. The increased spatial separation of the two molecular orbitals impedes their electronic exchange and thus explains the increased  $E_T$  of *o*-PCzTPD compared to *m*-PCzTPD. Thus, the *ortho* linkage effect<sup>33</sup> is further enhanced for these particular class of materials, owing to the steric demand of the TPD, which leads to an increased electronic separation of the molecular subunits, even compared to the corresponding *meta* derivatives.

### Electroluminescent properties

The title compounds were employed in PhOLED devices, to investigate their practical applicability as host materials. Owing to the higher  $E_T$  of the three materials compared to red emitting Ir(MDQ)<sub>2</sub>(acac) (2.0 eV),<sup>67</sup> Ir(MDQ)<sub>2</sub>(acac) was chosen as phosphorescent emitter for this purpose. PhOLED devices were fabricated with the structure of ITO/MoO<sub>3</sub> (8 nm)/TAPC (75 nm)/EML (12 nm)/BPhen (70 nm)/LiF/Al (Fig. S10, ESI†). The EMLs consisted of host materials *p*-PCzTPD (**R1**), *m*-PCzTPD (**R2**) or *o*-PCzTPD (**R3**) doped with Ir(MDQ)<sub>2</sub>(acac). A dopant concentration of 5% was applied without further optimization. TAPC was employed as hole transporting layer and BPhen was utilized as electron transporting and hole blocking layer. Current density–voltage–luminance and current efficiency–luminance–power efficiency curves of the three devices are depicted in Fig. 4 and the key electroluminescent properties are summarized in Table 3.

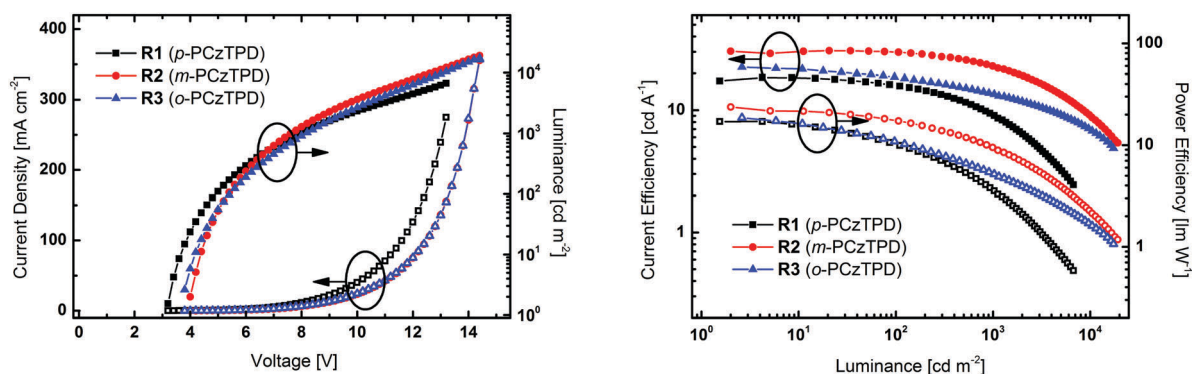
All devices exhibited electroluminescent emission exclusively from the Ir(MDQ)<sub>2</sub>(acac) dopant (Fig. S11, ESI†), indicating that exciton transfer from the host to the dopant or direct charge trapping on the phosphorescent emitter was effective. Moreover, all devices exhibited turn-on voltages between 3.2 V and 4.0 V. The slightly lower turn on voltage of 3.2 V of **R1** can be attributed to the lower bandgap and in particular the low lying LUMO level of *p*-PCzTPD compared to its regioisomers.

Among the three devices, *m*-PCzTPD based **R2** exhibited the best performance with a high maximum current efficiency (CE) of 30.8 cd A<sup>−1</sup> corresponding to a satisfying maximum external

**Table 2** Characteristic torsion angles  $\Phi_1$ – $\Phi_4$  of the least square planes of the aromatic ring systems of the materials under investigation

	$\Phi_1$ [°] (Cz1–Ph1)	$\Phi_2$ [°] (Ph1–TPD)	$\Phi_3$ [°] (TPD–Ph2)	$\Phi_4$ [°] (Ph2–Cz2)
<i>p</i> -PCzTPD <sup>a</sup>	54.53(12)	15.04(11)	18.08(11)	61.68(11)
<i>p</i> -PCzTPD <sup>b</sup>	55.67	19.14	20.30	55.52
<i>m</i> -CzTPD <sup>b</sup>	59.65	25.75	23.47	61.64
<i>o</i> -CzTPD <sup>b</sup>	67.16	49.43	53.46	67.91

<sup>a</sup> Determined by single-crystal X-ray diffraction. <sup>b</sup> Calculated applying density functional theory level (B3LYP/6-311+G\*).



**Fig. 4** Current density–voltage–luminance (left; luminance: full symbols, current density: hollow symbols) and current efficiency–luminance–power efficiency (right; current efficiency: full symbols, power efficiency: hollow symbols) curves of devices **R1**–**R3**.





Table 3 Electroluminescent properties of devices **R1–3**

	$V_{on}$ [V]	$CE^a$ [cd A <sup>-1</sup> ]	$PE^a$ [lm W <sup>-1</sup> ]	$EQE^a$ [%]
<b>R1</b>	3.2	15.9/9.1/18.5	10.1/3.6/17.1	10.1/5.8/11.8
<b>R2</b>	4.0	29.7/22.5/30.6	17.3/9.1/23.8	17.2/13.0/17.7
<b>R3</b>	3.8	18.4/13.6/22.5	10.7/5.2/18.6	10.5/7.7/12.9

<sup>a</sup> Measured at a brightness of 100 cd m<sup>-2</sup>/measured at a brightness of 1000 cd m<sup>-2</sup>/maximum efficiency.

quantum efficiency (EQE) of 17.7% and a maximum power efficiency (PE) of 23.8 lm W<sup>-1</sup>. Compared to **R2** devices **R1** and **R3** displayed somewhat lower efficiencies with a  $CE_{max}$  of 18.5 cd A<sup>-1</sup> and 22.2 cd A<sup>-1</sup> and a  $PE_{max}$  of 11.8 lm W<sup>-1</sup> and 12.9 lm W<sup>-1</sup>, respectively. The lower efficiency of **R1** may be attributed to the fact that the  $E_T$  of *p*-PCzTPD is rather close to the  $E_T$  of the red (MDQ)<sub>2</sub>(acac) dopant (2.0 eV),<sup>67</sup> which is also in accordance with the higher efficiency roll-off of **R1** compared to **R2** and **R3**. Although the performance of **R3** is slightly better compared to **R1**, it does not reach the efficiency values of **R2** despite a higher  $E_T$  of the host material. Nonetheless, the efficiency roll-off above a brightness of 1000 cd m<sup>-2</sup> is less pronounced in **R3**.

## Conclusions

In this study we incorporated the electron accepting thieno[3,4-*c*]pyrrole-4,6-dione building block in donor-acceptor host materials for PhOLEDs and prepared three isomeric derivatives for this purpose. The newly developed materials displayed favourable molecular properties for the application in red PhOLED devices and prototype devices employing Ir(MDQ)<sub>2</sub>(acac) as phosphorescent emitter featured satisfying current and power efficiencies. Thus, we have successfully demonstrated the applicability of the thieno[3,4-*c*]pyrrole-4,6-dione as acceptor in host materials for PhOLEDs and introduced this particular molecular unit as novel building block for functional organic materials for OLEDs.

## Experimental section

### General information

All reagents and solvents were purchased from commercial suppliers and used without further purification. Column chromatography was performed on silica 60 (Merck, 40–63 μm). NMR spectra were recorded on a Bruker Avance DRX-400 Spectrometer. A Thermo Scientific LTQ Orbitrap XL hybrid FTMS (Fourier Transform Mass Spectrometer) equipped with a Thermo Scientific MALDI source was used for high resolution mass spectrometry. The thermal analysis was carried out with a heating rate of 10 K min<sup>-1</sup> in a flowing argon atmosphere (25 mL min<sup>-1</sup>). TG measurements were performed on a Netzsch TG 209 F9 Tarsus system with open aluminium oxide crucibles. DSC measurements were performed on a Netzsch DSC 200 F3 Maia, working with aluminium pans with pierced lids. UV/Vis absorption and fluorescence emission spectra were recorded in DCM solutions (5 μM) with a Perkin Elmer Lambda 750 spectrometer and an Edinburgh FLS920, respectively. The time resolved low temperature phosphorescence spectra were

recorded in solid solutions (1 mg mL<sup>-1</sup>; toluene : EtOH = 9 : 1) at 77 K with a Perkin Elmer Instruments LS 50B luminance spectrometer. Cyclic voltammetry was performed using a three electrode configuration consisting of a Pt working electrode, a Pt counter electrode and an Ag/AgCl reference electrode and a PGSTAT128N, ADC164, DAC164, External, DI048 potentiostat provided by Metrohm Autolab B.V. Measurements were carried out in a 0.5 mM solution in anhydrous DCM with Bu<sub>4</sub>NBF<sub>4</sub> (0.1 M) as supporting electrolyte. The solutions were purged with nitrogen for 15 minutes prior to measurement. HOMO energy levels were calculated from the onset of the oxidation peaks, which was determined from the intersection of two tangents drawn at the background and the rising of the oxidation peaks.

### Synthetic details

General procedure for the synthesis of the host materials: Host materials *p*-PCzTPD, *m*-PCzTPD and *o*-PCzTPD were prepared using a Suzuki cross coupling reaction. A three necked flask was flushed with argon and charged with dibromide **1** (1.0 eq.) and boronic acid esters **2a–c** (2.5 eq.) before degassed THF (2.5 mM) and aqueous K<sub>2</sub>CO<sub>3</sub> (5.0 eq., 2 M) were added. Subsequently, Pd(PPh<sub>3</sub>)<sub>4</sub> (0.05 eq.) was added and the reaction mixture was heated to reflux until full conversion (TLC, 72 h). After cooling to room temperature the reaction mixture was poured on water and extracted with CH<sub>2</sub>Cl<sub>2</sub> three times. The combined organic layers were dried over Na<sub>2</sub>SO<sub>4</sub> and concentrated under reduced pressure. Purification of the crude product was accomplished by column chromatography.

**1,3-Bis[4-(9H-carbazol-9-yl)phenyl]-5-octyl-4H-thieno[3,4-*c*]pyrrole-4,6(5H)-dione (*p*-PCzTPD).** Starting from **1** (169 mg, 0.40 mmol), **2a** (370 mg, 1.00 mmol), K<sub>2</sub>CO<sub>3</sub> (277 mg, 2.00 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (23 mg, 0.02 mmol) *p*-PCzTPD was isolated after column chromatography (light petrol/toluene = 1/2) as yellow solid (199 mg, 0.27 mmol, 67%). <sup>1</sup>H NMR (400 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ = 8.46 (d, *J* = 8.5 Hz, 4H), 8.17 (d, *J* = 7.6 Hz, 4H), 7.76 (d, *J* = 8.5 Hz, 4H), 7.56 (d, *J* = 8.0 Hz, 4H), 7.46 (t, *J* = 7.6 Hz, 4H), 7.33 (t, *J* = 7.6 Hz, 4H), 3.71 (t, *J* = 7.0 Hz, 2H), 1.72 (q, *J* = 6.9 Hz, 2H), 1.44–1.24 (m, 10H), 0.88 (t, *J* = 6.9 Hz, 3H) ppm. <sup>13</sup>C NMR (100 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ = 163.4 (s), 144.3 (s), 141.0 (s), 139.9 (s), 131.7 (s), 130.3 (d), 129.9 (s), 127.6 (d), 126.7 (d), 124.2 (s), 121.0 (d), 120.9 (d), 110.4 (d), 39.2 (t), 32.4 (t), 29.8 (t), 29.8 (t), 29.0 (t), 27.5 (t), 23.2 (t), 14.4 (q) ppm. HRMS (MALDI): *m/z* calculated for C<sub>50</sub>H<sub>41</sub>N<sub>3</sub>O<sub>2</sub>S: 747.29140 [M]<sup>+</sup>, 748.29922 [M + H]<sup>+</sup>; found: 747.29229 [M]<sup>+</sup>, 748.29565 [M + H]<sup>+</sup>.

**1,3-Bis[4-(9H-carbazol-9-yl)phenyl]-5-octyl-4H-thieno[3,4-*c*]pyrrole-4,6(5H)-dione (*m*-PCzTPD).** Starting from **1** (412 mg, 0.97 mmol), **2b** (899 mg, 2.43 mmol), K<sub>2</sub>CO<sub>3</sub> (672 mg, 4.86 mmol) and Pd(PPh<sub>3</sub>)<sub>4</sub> (56 mg, 0.05 mmol) *m*-PCzTPD was isolated after column chromatography (light petrol/CH<sub>2</sub>Cl<sub>2</sub> = 3/1 → 1/1) as yellow solid (510 mg, 0.68 mmol, 70%). <sup>1</sup>H NMR (400 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ = 8.41 (s, 2H), 8.21–8.16 (m, 6H), 7.76–7.67 (m, 4H), 7.54 (d, *J* = 8.2 Hz, 4H), 7.45 (t, *J* = 7.3 Hz, 4H), 7.32 (t, *J* = 7.3 Hz, 4H), 3.63 (t, *J* = 7.2 Hz, 2H), 1.64 (q, *J* = 7.3 Hz, 2H), 1.36–1.19 (m, 10H), 0.85 (t, *J* = 6.7 Hz, 3H) ppm. <sup>13</sup>C NMR (100 MHz, CD<sub>2</sub>Cl<sub>2</sub>): δ = 163.3 (s), 144.3 (s), 141.2 (s), 139.0 (s), 132.9 (s), 132.1 (s), 131.1 (d), 129.0 (d), 127.5 (d), 127.1 (d), 126.7 (d), 124.1 (s), 120.8 (d),



120.8 (d), 110.4 (d), 39.2 (t), 32.4 (t), 29.7 (t), 29.7 (t), 28.9 (t), 27.5 (t), 23.2 (t), 14.4 (q) ppm. HRMS (MALDI):  $m/z$  calculated for  $C_{50}H_{41}N_3O_2S$ : 747.29140  $[M]^+$ , 748.29922  $[M + H]^+$ , 770.28117  $[M + Na]^+$ , found: 747.29282  $[M]^+$ , 748.29733  $[M + H]^+$ , 770.28271  $[M + Na]^+$ .

**1,3-Bis[2-(9H-carbazol-9-yl)phenyl]-5-octyl-4H-thieno[3,4-c]pyrrole-4,6(5H)-dione (o-PCzTPD).** Starting from **1** (566 mg, 1.34 mmol), **2c** (1233 mg, 3.31 mmol),  $K_2CO_3$  (922 mg, 6.67 mmol) and  $Pd(PPh_3)_4$  (77 mg, 0.07 mmol) **o-PCzTPD** was isolated after column chromatography (light petrol/ $CH_2Cl_2$  = 1/1) as yellow solid (172 mg, 0.23 mmol, 17%).  $^1H$  NMR (400 MHz,  $CD_2Cl_2$ ):  $\delta$  = 7.96 (d,  $J$  = 7.2 Hz, 4H), 7.87–7.45 (m, 2H), 7.53–7.48 (m, 4H), 7.26–7.24 (m, 2H), 7.19–7.11 (m, 8H), 6.75 (d,  $J$  = 7.6 Hz, 4H), 3.61 (t,  $J$  = 7.3 Hz, 2H) 1.55 (q,  $J$  = 7.2 Hz, 2H), 1.37–1.21 (m, 10H), 0.91 (t,  $J$  = 6.7 Hz, 3H) ppm.  $^{13}C$  NMR (100 MHz,  $CD_2Cl_2$ ):  $\delta$  = 162.8 (s), 141.8 (s), 141.7 (s), 135.9 (s), 132.6 (d), 132.0 (s), 131.8 (d), 129.9 (d), 129.8 (s), 128.9 (d), 126.4 (d), 124.0 (s), 120.6 (d), 120.6 (d), 110.2 (d), 38.7 (t), 32.4 (t), 29.8 (t), 29.8 (t), 28.8 (t), 27.4 (t), 23.2 (t), 14.5 (q) ppm. HRMS (MALDI):  $m/z$  calculated for  $C_{50}H_{41}N_3O_2S$ : 747.29140  $[M]^+$ , 748.29922  $[M + H]^+$ , 770.28117  $[M + Na]^+$ , found: 747.29249  $[M]^+$ , 748.29516  $[M + H]^+$ , 770.28228  $[M + Na]^+$ .

### Computational details

The (TD)DFT computations were performed using the Gaussian 09 package, revision D.01.<sup>68</sup> Density functional theory (DFT) and time-dependent (TD)DFT calculations were performed using the Becke three parameters hybrid functional with Lee–Yang–Perdew correlation (B3LYP),<sup>69,70</sup> in combination with Pople basis sets (6-31G\*, 6-311+G\*).<sup>71</sup> The geometry optimizations were performed in gas phase and without symmetry constraints. For the calculation of HOMO/LUMO levels, the ground state ( $S_0$ ) geometries were optimized applying the 6-311+G\* basis set. The determination of the triplet energy was achieved by the calculation of the  $T_1$  excitation energy applying TDDFT level and the 6-311+G\* basis to a  $S_0$  geometry optimized at DFT level using the 6-31G\* basis set. Orbital plots were generated using GaussView.<sup>72</sup>

### Single crystal diffraction

A crystal of **p-PCzTPD** suitable for single crystal diffraction was selected under a polarizing microscope, embedded in perfluorinated oil and attached to Kapton<sup>®</sup> micromounts. Intensity data were collected in a dry stream of nitrogen at 100 K on a Bruker KAPPA APEX II diffractometer system. Data were reduced using SAINT-Plus<sup>73</sup> and an empirical absorption correction using the multi-scan approach implemented in SADABS<sup>73</sup> was applied. The crystal structures were solved by charge-flipping implemented in SUPERFLIP<sup>74</sup> and refined against  $F$  with the JANA2006<sup>75</sup> software package. The non-H atoms were refined with anisotropic displacement parameters. The H atoms were placed at calculated positions and refined as riding on the parent C atoms.

### Device fabrication and measurement

All synthesized materials were additionally purified by crystallization (light petrol with minor amounts of toluene) prior to device fabrication. OLED fabrication and testing: the fabricated

devices were grown on clean glass substrates pre-coated with ITO with a sheet resistance of  $10 \Omega \text{ sq}^{-1}$ . After cleaning with detergent and deionized water, they were dried in an oven at  $120^\circ\text{C}$  for 30 minutes and then treated with UV-ozone for 15 minutes. All layers were grown in succession by thermal evaporation without breaking the vacuum ( $\sim 10^{-4}$  Pa). The evaporation rates for the organic layer were in a range of  $1\text{--}2 \text{ \AA s}^{-1}$ , LiF and Al were evaporated with a rate of approximately  $0.1$  and  $5.0 \text{ \AA s}^{-1}$ , respectively. All measurements were carried out in ambient atmosphere. The current–voltage–brightness characteristics were recorded using a Keithley source measurement unit (Keithley 2400 and Keithley 2000) with a calibrated silicon photodiode. The electroluminescence (EL) spectra were measured with a Spectrascan PR650 spectrophotometer.

## Acknowledgements

This work was supported in part by the TU Wien “Innovative Projects” research funds and the Austrian Federal Ministry of Science, Research and Economy. The X-ray centre of the TU Wien is acknowledged for providing access to the single-crystal diffractometer. P. K. and J. F. gratefully acknowledge financial support by the Austrian Science Fund (FWF) (Grant No. I 2589-N34). J. C. and D. M. gratefully acknowledge financial support by the National Natural Science Foundation of China (Grant No. 11661131001).

## Notes and references

- M. Pope, P. Magnante and H. P. Kallmann, *J. Chem. Phys.*, 1963, **38**, 2042–2043.
- W. Helfrich and W. G. Schneider, *Phys. Rev. Lett.*, 1965, **14**, 229–231.
- C. W. Tang and S. A. VanSlyke, *Appl. Phys. Lett.*, 1987, **51**, 913–915.
- J. H. Burroughes, D. D. C. Bradley, A. R. Brown, R. N. Marks, K. Mackay, R. H. Friend, P. L. Burn and A. B. Holmes, *Nature*, 1990, **347**, 539–541.
- D. Braun and A. J. Heeger, *Appl. Phys. Lett.*, 1991, **58**, 1982–1984.
- S. R. Forrest, *Nature*, 2004, **428**, 911–918.
- M. C. Gather, A. Köhnen and K. Meerholz, *Adv. Mater.*, 2011, **23**, 233–248.
- S. Reineke, M. Thomschke, B. Lüssem and K. Leo, *Rev. Mod. Phys.*, 2013, **85**, 1245–1293.
- M. A. Baldo, D. F. O'Brien, Y. You, A. Shoustikov, S. Sibley, M. E. Thompson and S. R. Forrest, *Nature*, 1998, **395**, 151–154.
- M. A. Baldo, S. Lamansky, P. E. Burrows, M. E. Thompson and S. R. Forrest, *Appl. Phys. Lett.*, 1999, **75**, 4–6.
- H. Uoyama, K. Goushi, K. Shizu, H. Nomura and C. Adachi, *Nature*, 2012, **492**, 234–238.
- Y. Sun, N. C. Giebink, H. Kanno, B. Ma, M. E. Thompson and S. R. Forrest, *Nature*, 2006, **440**, 908–912.
- S. Reineke, F. Lindner, G. Schwartz, N. Seidler, K. Walzer, B. Lüssem and K. Leo, *Nature*, 2009, **459**, 234–238.
- Y. Tao, K. Yuan, T. Chen, P. Xu, H. H. Li, R. F. Chen, C. Zheng, L. Zhang and W. Huang, *Adv. Mater.*, 2014, **26**, 7931–7958.



- 15 Q. Zhang, B. Li, S. Huang, H. Nomura, H. Tanaka and C. Adachi, *Nat. Photonics*, 2014, **8**, 326–332.
- 16 W.-C. Chen, C.-S. Lee and Q.-X. Tong, *J. Mater. Chem. C*, 2015, **3**, 10957–10963.
- 17 R. J. Holmes, S. R. Forrest, Y.-J. Tung, R. C. Kwong, J. J. Brown, S. Garon and M. E. Thompson, *Appl. Phys. Lett.*, 2003, **82**, 2422–2424.
- 18 S. Tokito, T. Iijima, Y. Suzuri, H. Kita, T. Tsuzuki and F. Sato, *Appl. Phys. Lett.*, 2003, **83**, 569–571.
- 19 L. Xiao, Z. Chen, B. Qu, J. Luo, S. Kong, Q. Gong and J. Kido, *Adv. Mater.*, 2011, **23**, 926–952.
- 20 K. S. Yook and J. Y. Lee, *Adv. Mater.*, 2012, **24**, 3169–3190.
- 21 H. Sasabe and J. Kido, *Eur. J. Org. Chem.*, 2013, 7653–7663.
- 22 Y. Tao, C. Yang and J. Qin, *Chem. Soc. Rev.*, 2011, **40**, 2943–2970.
- 23 A. Chaskar, H.-F. Chen and K.-T. Wong, *Adv. Mater.*, 2011, **23**, 3876–3895.
- 24 S. O. Jeon and J. Y. Lee, *J. Mater. Chem.*, 2012, **22**, 4233–4243.
- 25 B. E. Koene, D. E. Loy and M. E. Thompson, *Chem. Mater.*, 1998, **10**, 2235–2250.
- 26 Z. Ge, T. Hayakawa, S. Ando, M. Ueda, T. Akiike, H. Miyamoto, T. Kajita and M.-a. Kakimoto, *Adv. Funct. Mater.*, 2008, **18**, 584–590.
- 27 Y. Tao, Q. Wang, Y. Shang, C. Yang, L. Ao, J. Qin, D. Ma and Z. Shuai, *Chem. Commun.*, 2009, 77–79.
- 28 C. H. Chen, W. S. Huang, M. Y. Lai, W. C. Tsao, J. T. Lin, Y. H. Wu, T. H. Ke, L. Y. Chen and C. C. Wu, *Adv. Funct. Mater.*, 2009, **19**, 2661–2670.
- 29 Y. Tao, Q. Wang, L. Ao, C. Zhong, C. Yang, J. Qin and D. Ma, *J. Phys. Chem. C*, 2010, **114**, 601–609.
- 30 Y. Kuwabara, H. Ogawa, H. Inada, N. Noma and Y. Shirota, *Adv. Mater.*, 1994, **6**, 677–679.
- 31 Z. Ge, T. Hayakawa, S. Ando, M. Ueda, T. Akiike, H. Miyamoto, T. Kajita and M.-a. Kakimoto, *Org. Lett.*, 2008, **10**, 421–424.
- 32 S.-J. Su, H. Sasabe, T. Takeda and J. Kido, *Chem. Mater.*, 2008, **20**, 1691–1693.
- 33 Y. Tao, Q. Wang, C. Yang, C. Zhong, K. Zhang, J. Qin and D. Ma, *Adv. Funct. Mater.*, 2010, **20**, 304–311.
- 34 H.-C. Ting, Y.-M. Chen, H.-W. You, W.-Y. Hung, S.-H. Lin, A. Chaskar, S.-H. Chou, Y. Chi, R.-H. Liu and K.-T. Wong, *J. Mater. Chem.*, 2012, **22**, 8399–8407.
- 35 P. Kautny, D. Lumpi, Y. Wang, A. Tissot, J. Bintliger, E. Horkel, B. Stöger, C. Hametner, H. Hagemann, D. Ma and J. Fröhlich, *J. Mater. Chem. C*, 2014, **2**, 2069–2081.
- 36 D. Zhang, L. Duan, C. Li, Y. Li, H. Li, D. Zhang and Y. Qiu, *Adv. Mater.*, 2014, **26**, 5050–5055.
- 37 P. Kautny, Z. Wu, J. Eichelner, E. Horkel, B. Stöger, J. Chen, D. Ma, J. Fröhlich and D. Lumpi, *Org. Electron.*, 2016, **34**, 237–245.
- 38 J. Lee, K. Shizu, H. Tanaka, H. Nomura, T. Yasuda and C. Adachi, *J. Mater. Chem. C*, 2013, **1**, 4599–4604.
- 39 H. Tanaka, K. Shizu, H. Nakanotani and C. Adachi, *Chem. Mater.*, 2013, **25**, 3766–3771.
- 40 S. Y. Lee, T. Yasuda, Y. S. Yang, Q. Zhang and C. Adachi, *Angew. Chem., Int. Ed.*, 2014, **126**, 6520–6524.
- 41 Q. Zhang, H. Kuwabara, W. J. Potscavage, S. Huang, Y. Hatae, T. Shibata and C. Adachi, *J. Am. Chem. Soc.*, 2014, **136**, 18070–18081.
- 42 Y. Hamada, C. Adachi, T. Tsutsui and S. Saito, *Jpn. J. Appl. Phys., Part 1*, 1992, **31**, 1812–1816.
- 43 C. Adachi, T. Tsutsui and S. Saito, *Appl. Phys. Lett.*, 1989, **55**, 1489–1491.
- 44 P. Kautny, Z. Wu, B. Stöger, A. Tissot, E. Horkel, J. Chen, D. Ma, H. Hagemann, J. Fröhlich and D. Lumpi, *Org. Electron.*, 2015, **17**, 216–228.
- 45 D. F. O'Brien, M. A. Baldo, M. E. Thompson and S. R. Forrest, *Appl. Phys. Lett.*, 1999, **74**, 442–444.
- 46 S.-J. Su, Y. Takahashi, T. Chiba, T. Takeda and J. Kido, *Adv. Funct. Mater.*, 2009, **19**, 1260–1267.
- 47 Z. Gao, C. S. Lee, I. Bello, S. T. Lee, R.-M. Chen, T.-Y. Luh, J. Shi and C. W. Tang, *Appl. Phys. Lett.*, 1999, **74**, 865–867.
- 48 H. Inomata, K. Goushi, T. Masuko, T. Konno, T. Imai, H. Sasabe, J. J. Brown and C. Adachi, *Chem. Mater.*, 2004, **16**, 1285–1291.
- 49 D. R. Lee, M. Kim, S. K. Jeon, S.-H. Hwang, C. W. Lee and J. Y. Lee, *Adv. Mater.*, 2015, **27**, 5861–5867.
- 50 J. Kido, C. Ohtaki, K. Hongawa, K. Okuyama and K. Nagai, *Jpn. J. Appl. Phys., Part 2*, 1993, **32**, L917–L920.
- 51 P. E. Burrows, A. B. Padmaperuma, L. S. Sapochak, P. Djurovich and M. E. Thompson, *Appl. Phys. Lett.*, 2006, **88**, 183503.
- 52 S. O. Jeon, K. S. Yook, C. W. Joo and J. Y. Lee, *Adv. Funct. Mater.*, 2009, **19**, 3644–3649.
- 53 C. Han, G. Xie, H. Xu, Z. Zhang, L. Xie, Y. Zhao, S. Liu and W. Huang, *Adv. Mater.*, 2011, **23**, 2491–2496.
- 54 H. Wang, L. Xie, Q. Peng, L. Meng, Y. Wang, Y. Yi and P. Wang, *Adv. Mater.*, 2014, **26**, 5198–5204.
- 55 C. W. Lee and J. Y. Lee, *Adv. Mater.*, 2013, **25**, 5450–5454.
- 56 Y. Im and J. Y. Lee, *Chem. Commun.*, 2013, **49**, 5948–5950.
- 57 T. Nakagawa, S.-Y. Ku, K.-T. Wong and C. Adachi, *Chem. Commun.*, 2012, **48**, 9580–9582.
- 58 C. Piliego, T. W. Holcombe, J. D. Douglas, C. H. Woo, P. M. Beaujuge and J. M. J. Fréchet, *J. Am. Chem. Soc.*, 2010, **132**, 7595–7597.
- 59 T.-Y. Chu, J. Lu, S. Beaupré, Y. Zhang, J.-R. Pouliot, S. Wakim, J. Zhou, M. Leclerc, Z. Li, J. Ding and Y. Tao, *J. Am. Chem. Soc.*, 2011, **133**, 4250–4253.
- 60 C. B. Nielsen, R. S. Ashraf, B. C. Schroeder, P. D'Angelo, S. E. Watkins, K. Song, T. D. Anthopoulos and I. McCulloch, *Chem. Commun.*, 2012, **48**, 5832–5834.
- 61 M.-C. Yuan, M.-Y. Chiu, S.-P. Liu, C.-M. Chen and K.-H. Wei, *Macromolecules*, 2010, **43**, 6936–6938.
- 62 X. Guo, N. Zhou, S. J. Lou, J. Smith, D. B. Tice, J. W. Hennek, R. P. Ortiz, J. T. L. Navarrete, S. Li, J. Strzalka, L. X. Chen, R. P. H. Chang, A. Facchetti and T. J. Marks, *Nat. Photonics*, 2013, **7**, 825–833.
- 63 X. Guo, R. P. Ortiz, Y. Zheng, M.-G. Kim, S. Zhang, Y. Hu, G. Lu, A. Facchetti and T. J. Marks, *J. Am. Chem. Soc.*, 2011, **133**, 13685–13697.
- 64 H. Li, Y. Wang, K. Yuan, Y. Tao, R. Chen, C. Zheng, X. Zhou, J. Li and W. Huang, *Chem. Commun.*, 2014, **50**, 15760–15763.
- 65 Y. Tao, Q. Wang, L. Ao, C. Zhong, J. Qin, C. Yang and D. Ma, *J. Mater. Chem.*, 2010, **20**, 1759–1765.
- 66 E. Mondal, W.-Y. Hung, H.-C. Dai and K.-T. Wong, *Adv. Funct. Mater.*, 2013, **23**, 3096–3105.



- 67 G. Schwartz, S. Reineke, T. Rosenow, K. Walzer and K. Leo, *Adv. Funct. Mater.*, 2009, **19**, 1319–1333.
- 68 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery Jr., J. E. Peralta, F. Ogliaro, M. J. Bearpark, J. Heyd, E. N. Brothers, K. N. Kudin, V. N. Staroverov, R. Kobayashi, J. Normand, K. Raghavachari, A. P. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, N. J. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, Ö. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski and D. J. Fox, *Gaussian 09, Revision D.01*, Gaussian, Inc., Wallingford, CT, USA, 2009.
- 69 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785–789.
- 70 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 71 R. Krishnan, J. S. Binkley, R. Seeger and J. A. Pople, *J. Chem. Phys.*, 1980, **72**, 650–654.
- 72 R. Dennington, T. Keith and J. Millam, *GaussView, Version 5*, Semichem, Inc., Shawnee Mission, KS, 2009.
- 73 *Bruker computer programs: APEX2, SAINT and SADABS*, Bruker AXS Inc., Madison, WI, 2015.
- 74 L. Palatinus and G. Chapuis, *J. Appl. Crystallogr.*, 2007, **40**, 786–790.
- 75 V. Petříček, M. Dušek and L. Palatinus, *Z. Kristallogr. – Cryst. Mater.*, 2014, **229**, 345–352.

