

# Effect of Polymerization Statistics on the Electronic Properties of Copolymers for Organic Photovoltaics

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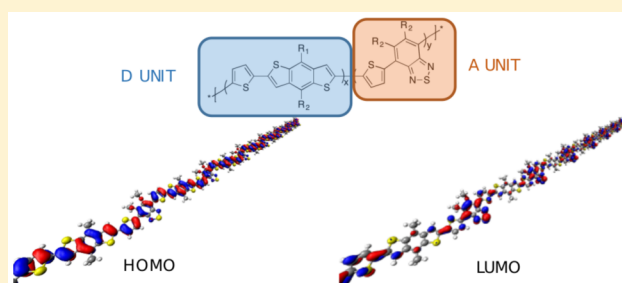
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## S Supporting Information

**ABSTRACT:** Statistical block copolymers, composed of donor (D) and acceptor (A) blocks, are a novel type of material for organic photovoltaics (OPVs) devices. In particular a new series of polymers based on PBTZT-*stat*-BDTT-8, recently developed by Merck, offers high solubility in different solvents, and a high power conversion efficiency (PCE) in different device architectures. Although it is known that the electronic properties of these materials may be significantly affected by attaching different functional groups on different blocks, it is not fully clear how important the influence of the polymer composition (i.e., the D/A block ratio) is, even if previous studies suggest that this might also have an effect. Therefore, the effect of the polymer composition in terms of HOMO, LUMO energies, and band gap was explored by studying a number of long chain oligomers with more than 1000 atoms each and with different D/A ratios. This study, that is novel both conceptually and methodologically, was made possible by using the linear-scaling reformulation of DFT implemented in the ONETEP code. Our results showed that changing the composition has a significant effect on the electronic structure of statistical copolymers, making this an alternative and suitable strategy to obtain materials with desired properties. Also, a systematic analysis of the effect of a range of different substituents placed in the D and A blocks of PBTZT-*stat*-BDTT-8 was performed in order to investigate how this class of materials responds to functionalization. We found that it is not possible to know a priori using chemical intuition what kind of influence different types of functional groups may have on these systems, and therefore, computational modeling is essential.



## INTRODUCTION

The world energy consumption is constantly growing, and therefore, there is a pressing need to find new and alternative sources of energy in both industry and academia. Technologies based on renewable and clean energies do exist, and among these, photovoltaics show great promise. Organic photovoltaics (OPV) based on  $\pi$ -conjugated polymers, in particular, have drawn attention in recent years because of their low manufacturing costs, solution processability, and flexibility, not to mention the fact that these devices are lightweight and also efficient in diffuse light and indoors.<sup>1</sup> Moreover, despite their relatively low performance, they show a very fast energy payback time (EPBT): while for silicon-based technologies the EPBT is in the order of years, for OPV it has been estimated to be in the order of days.<sup>2,3</sup>

The active layer of an OPV module, in the state-of-the-art bulk heterojunction (BHJ) architecture,<sup>4</sup> consists of a blend of electron-donor and electron-acceptor materials, the first being typically a conjugated polymer and the latter being in the

majority of cases a soluble fullerene derivative ([6,6]-phenyl-C<sub>61</sub>-butyric acid methyl ester, also known as PC<sub>61</sub>BM or simply PCBM). This kind of interpenetrating network allows the exciton splitting and the subsequent generation of an electron and a hole in the form of free charge carriers. Since the photoabsorption happens in the polymer bulk, the photo-current conversion efficiency (PCE) largely depends on the number of photons that are absorbed by the polymer. One of the most used materials in BHJ devices was poly(3-hexylthiophene) (P3HT), a regioregular homopolymer that led to modules showing a PCE of 6%.<sup>5,6</sup> One of the problems of P3HT, however, lied in its relatively large band gap<sup>7</sup> (~2 eV) that does not allow the absorption of a large number of photons, since its absorption spectrum overlaps with just a small part of the solar emission spectrum; it has in fact been

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estimated that P3HT is able to collect in the best case scenario only ~20% of the solar photons.<sup>8</sup> Different strategies were then proposed with the aim of lowering the band gap of these materials, in order to harvest photons at longer wavelengths and therefore absorb a wider portion of the solar radiation that reaches the Earth:<sup>9</sup> one of these is the synthesis of donor–acceptor (D–A or push–pull) copolymers, formed by alternating electron-rich and electron-deficient blocks. This leads to an intramolecular charge transfer (ICT) process between the D and A blocks,<sup>10</sup> and to materials having, in general, a smaller and tunable band gap with respect to homopolymers like P3HT mentioned above; this effect is due to the orbital mixing between the two different units. The band gap (and, consequently, HOMO and LUMO energies) can be carefully tuned by combining different acceptor and donor units and by changing the functional groups attached to those blocks. In addition, recent studies suggest that modifying the composition (i.e., the donor–acceptor ratio) of the material can lead to different HOMO and LUMO energies.<sup>11</sup> This finding can potentially offer a novel strategy for the design of polymers with desired electronic properties. In general, it is desirable to reduce the band gap while maintaining at the same time a low HOMO level so that a high open-circuit voltage ( $V_{OC}$ ) value, and in turn, a high efficiency can be obtained, as proposed by Scharber and co-workers.<sup>12</sup> It has in fact been demonstrated empirically that for BHJ solar cells with PCBM as the acceptor species, an approximate linear relation exists between the HOMO energy of the conjugated polymer and the  $V_{OC}$  of the device:

$$V_{OC} = (1/e)(|E_{HOMO}^{donor}| - |E_{LUMO}^{PCBM}|) - 0.3V \quad (1)$$

where  $e$  is the elementary charge, the LUMO energy of PCBM is taken as a constant, and 0.3V is an empirical value. Moreover, ideally, in order to have a good hole mobility throughout a copolymer chain, valence bands (in other words, HOMO energies) of different blocks should be similar.

One of the top D–A polymers in terms of efficiency, solubility, and solution processability is part of the PBTZT-*stat*-BDTT-8 series, developed by Merck and employed in the solar tree installation at the German pavilion in Milan EXPO 2015.<sup>13</sup> This royal blue polymer, with a band gap of 1.7 eV, shows a 9% PCE in lab scale and 4.5% PCE in a fully roll-to-roll (R2R) module, the latter being the highest reported performance of flexible R2R semitransparent modules. PBTZT-*stat*-BDTT-8 is a novel class of OPV materials, consisting of statistical block copolymers composed of substituted benzo-[1,2-*b*:4,5-*b'*]dithiophene (BDT), thiophene (T), and 2,1,3-benzothiadiazole (BTZ). The fact that this material is a statistical copolymer means that blocks may have different lengths, different electronic properties, and may not be present with a 1:1 ratio in the chain: in fact, it has been observed that these materials behave differently with respect to regioregular analogues.<sup>14</sup> In addition, due to their reduced regularity, they show a higher solubility in nonhalogenated solvents, and this is of great importance from an industrial perspective, since widely used solvents such as chlorobenzene (CB) or *ortho*-dichlorobenzene (*o*-DCB) are not suitable for mass production.<sup>15</sup>

The main focus of this paper is the effect of composition, that is, the effect of different block ratios in the copolymer chain, in terms of HOMO, LUMO energies, and band gap, by studying a set of ~1000-atom oligomers, each one with a different D/A block ratio. This study, which is both conceptually and

methodologically novel, was enabled by using a linear-scaling DFT approach that allowed the investigation of the electronic properties of these systems at a far larger scale than possible before. We have also studied the effect of a range of different substituents both in the donor and acceptor units of some analogues of PBTZT-*stat*-BDTT-8, to investigate how and in what measure the electronic properties of this class of statistical copolymers can be affected by replacing the side chains in the D and A blocks with electron-donating and electron-withdrawing functional groups.

This study has been carried out using density functional theory calculations. The methods that have been used are described in [Methods](#), including the setup details of the simulations, while in [Results and Discussion](#), results are presented, together with their interpretation in the context of OPVs. Finally, conclusions are discussed in [Conclusions](#).

## METHODS

**The ONETEP Program.** All simulations were performed in the density functional theory (DFT) framework of electronic structure theory using the ONETEP (Order-N Electronic Total Energy Package) code for linear-scaling DFT.<sup>16,17</sup> Linear-scaling reformulations of DFT have been known, but suffer from reduced accuracy (i.e., only work with small basis sets); the peculiarity of ONETEP is that it combines linear-scaling behavior with large basis set accuracy, allowing efficient calculations also on systems composed of thousands of atoms. To achieve linear-scaling behavior, DFT has been reformulated in terms of the one-particle density matrix. In terms of Kohn–Sham orbitals, the density matrix can be expressed as follows

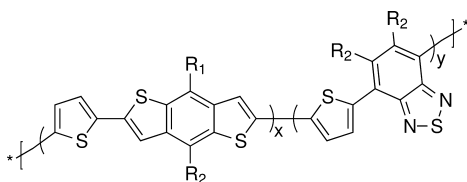
$$\rho(\mathbf{r}, \mathbf{r}') = \sum_{n=0}^{\infty} f_n \psi_n(\mathbf{r}) \psi_n^*(\mathbf{r}') \quad (2)$$

where  $f_n$  is the occupancy and  $\psi_n(\mathbf{r})$  are the Kohn–Sham orbitals. In ONETEP, the density matrix assumes the form

$$\rho(\mathbf{r}, \mathbf{r}') = \sum_{\alpha} \sum_{\beta} \phi_{\alpha}(\mathbf{r}) K^{\alpha\beta} \phi_{\beta}^*(\mathbf{r}') \quad (3)$$

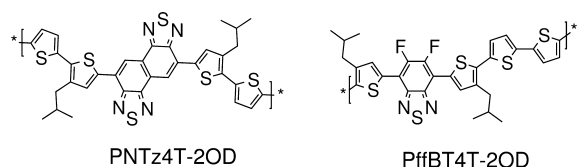
where  $\phi_{\alpha}(\mathbf{r})$  are local orbitals called nonorthogonal generalized Wannier functions (NGWFs)<sup>18</sup> and  $K$  is known as the density kernel. Making the density kernel sparse by truncation and enforcing strict localization of the NGWFs onto atomic regions leads to linear-scaling behavior. To achieve plane-wave accuracy, both the density kernel and the NGWFs, that are confined into spherical regions centered on each one of the atoms, are optimized during the calculation. The NGWFs are expanded in a basis set of periodic sinc (psinc) functions<sup>19</sup> equivalent to a plane-wave basis, allowing to easily control the basis set quality with just one parameter, equivalent to the kinetic energy cutoff in conventional DFT codes that make use of plane waves.

**Simulation Details.** Three limiting cases of structures were considered:  $x = 0$ ,  $y = 1$  and  $x = 1$ ,  $x = 0$ , where  $x$  and  $y$  are the number of units for each block, BDT-T and BTZ-T, respectively, as shown in [Figure 1](#); in other words, each single block alone was studied, as well as structures with alternating electron-rich and electron-deficient units. The effect of different substituents was then explored, in particular:  $R_1$ ,  $R_2 = H$ , F,  $OCH_3$ ,  $CH_3$ , CN; in a few cases, the  $OCH_3$  and  $CH_3$  groups were also replaced with the bulkier  $OC(CH_3)_3$  and  $C(CH_3)_3$  groups, respectively, in order to test the effect of steric



**Figure 1.** Structure of PBTZT-*stat*-BDTT-8, with  $x$  being the BDT-T block (donor, D) and  $y$  the BTZ-T block (acceptor, A).

hindrance on the electronic structure of these materials. The effect of fluorination on thiophene rings was also considered, as well as the replacement of benzothiadiazole with naphtho[1,2-*c*:5,6-*c'*]bis[1,2,5]thiadiazole (NTZ). As comparative examples, two high-performing polymers were also taken into consideration: PffBT4T-2OD<sup>20</sup> and PNT4T-2OD<sup>21</sup> (see Figure 2).



**Figure 2.** Structures of PNTz4T-2OD and PffBT4T-2OD, two examples of high-performing polymers.

The electronic structures of both repeat units and polymers were studied, with a specific focus on the HOMO and LUMO levels, as well as the band gap ( $E_g$ ); the structures of polymers were obtained using periodic boundary conditions (PBC) as implemented in ONETEP. When needed, in order to converge band gaps, more than a single unit was used to build the repeating cells. Note that for structures with  $x = 0$  and  $y = 0$  (i.e., BTZ-T, A only, and BDT-T, D only) a single unit consisted of two alternated benzothiadiazole and thiophene moieties, and two alternated benzodithiophene and thiophene units, respectively. It is worth mentioning that besides PBC, other methods to predict band gaps of conjugated polymers have been reported in literature, such as the extrapolation of the linear (or quadratic) curve of the band gap against  $1/n$ , where  $n$  is the number of monomer units.<sup>22</sup> A comparative study between those methods for a subset of structures studied in this paper is available in the [Supporting Information](#).

Finally, to overcome the limitation of being able to treat only perfectly alternating copolymers and to allow the study of polymer chains with different block ratios, five oligomers, each one composed of a total sum of 40 D and A blocks, consisting of  $\sim 1000$  atoms were built using the Materials Studio<sup>23</sup> software by setting different values of reactivity ratios for each block; their geometries were then optimized and their electronic structures were analyzed. The reactivity ratios used in this work refer to the Mayo–Lewis equation<sup>24</sup> (known also as copolymer equation) and they describe whether a monomer A is more likely to react with itself rather than with a monomer B during a copolymerization reaction; if the reactivity ratios are known, then it is possible to predict the final composition of the resulting copolymer. Note that every oligomer presents the same functional groups in  $R_1$  and  $R_2$  positions, that is  $\text{CH}_3$  and  $\text{OCH}_3$ , respectively. In order to avoid the effect of chain twisting and to make all results comparable, the starting guess structure for each oligomer before geometry optimization was a flat and linear conformation. Table 1 shows the reactivity ratios of the D and A units used for each oligomer, together with the

number of D and A blocks that compose each oligomer, and the total number of atoms.

**Table 1.** Reactivity Ratios Used in This Work for BDT-T (D unit) and BTZ-T (A unit) for Each Studied Oligomer, Together with the Number of D and A Blocks Present in Each Chain, and the Total Number of Atoms

$r_D$	$r_A$	no. D blocks	no. A blocks	no. atoms
10	0.1	38	2	1156
5	0.2	36	4	1150
0.2	5	9	31	1069
0.1	10	4	36	1054
1	1	18	22	1096

All the ground-state energy calculations were done in vacuum using the Perdew–Burke–Ernzerhof exchange–correlation functional (PBE)<sup>25</sup> with the projector augmented wave method (PAW). It is well-known that despite giving a good description of the properties of a chemical system, pure functionals, such as PBE, fail to accurately describe the  $E_g$  of semiconductors: the width of the gap is underestimated.<sup>26,27</sup> A better approximation of the  $E_g$  can be obtained with hybrid exchange–correlation functionals such as B3LYP.<sup>28,29</sup> Since the available ONETEP version allows for periodic calculations using nonhybrid functionals only,<sup>30</sup> the hybrid functional B3LYP was also used together with norm-conserving pseudopotentials only when running simulations on repeat units. After calibration of the method, optimal values for the kinetic energy cutoff and the NGWFs radii were found to be respectively 800 eV and 9.0 bohr, and they were used for all the simulations.

## RESULTS AND DISCUSSION

**Electronic Levels and Band Gaps.** For each studied structure (see Figures 1 and 2), detailed results of the energy levels (HOMO, LUMO, and  $E_g$ ) are shown in Table 2 both for repeat units and polymers, and in Figures 3 and 4 for repeat units and polymers, respectively. Structures from 1 to 12 all consist of alternating D–A blocks (note that in each structure the A unit corresponds to BTZ-T, with the only exception of 12 in which the BTZ-T moiety has been substituted with NTZ-T), structures from 13 to 16 consist of only A blocks (BTZ-T), while the ones from 17 to 21 are composed of just D blocks (BDT-T). Lastly, structures 22 and 23 are PNT4T-2OD and PffBT4T-2OD, respectively, two high-performing polymers studied in literature, as shown in Figure 2.

Results show that on average, both for repeat units and polymers, structures 13–16 composed of electron-deficient BTZ-T (A unit) present smaller gaps with respect to structures 17–21 composed of electron-rich BDT-T (D unit); structures 1–12, obtained by combining the previously discussed A and D units, present intermediate values as expected. In addition it can be seen that electron-withdrawing groups like F and CN tend to stabilize, and therefore shift to lower values, both HOMO and LUMO levels in both A and D units, while the opposite effect is observed for electron-donating groups like  $\text{CH}_3$  and  $\text{OCH}_3$ .

Focusing on the energy gaps of repeat units, it is quite remarkable that not only the values obtained using the B3LYP functional follow the same trend of the ones obtained with PBE, but also that their absolute difference is rather constant, and close to 1 eV, as shown in Figure 3. The same behavior can be observed for HOMO and LUMO energies: the effect of

Table 2. HOMO, LUMO, and  $E_g$  Obtained with ONETEP for All the Repeat Units and Polymer Structures Studied in This Paper<sup>a</sup>

#	$R_1/R_2$	repeat units						polymers		
		energies (eV); PBE			energies (eV); B3LYP			energies (eV); PBE		
		HOMO	LUMO	$E_g$	HOMO	LUMO	$E_g$	HOMO	LUMO	$E_g$
1	H/H	-4.75	-3.39	1.36	-5.15	-2.79	2.36	-4.13	-3.22	0.91
2	CH <sub>3</sub> /H	-4.66	-3.36	1.30	-5.06	-2.75	2.32	-4.04	-3.17	0.87
3	H/OCH <sub>3</sub>	-4.66	-3.23	1.43	-5.06	-2.62	2.44	-4.06	-3.05	1.01
4	CH <sub>3</sub> /OCH <sub>3</sub>	-4.54	-3.17	1.37	-4.94	-2.56	2.38	-4.20	-3.25	0.95
5	CH <sub>3</sub> /OC(CH <sub>3</sub> ) <sub>3</sub>	-4.51	-3.13	1.38	-4.91	-2.51	2.40	-3.92	-2.95	0.97
6	OCH <sub>3</sub> /OCH <sub>3</sub>	-4.18	-3.14	1.04	-4.68	-2.55	2.13	-3.75	-2.98	0.77
7	C(CH <sub>3</sub> ) <sub>3</sub> /OCH <sub>3</sub>	-4.50	-3.17	1.33	-4.91	-2.56	2.35	-3.94	-2.98	0.96
8	CN/OCH <sub>3</sub>	-5.03	-3.56	1.47	-5.43	-2.97	2.46	-4.84	-3.80	1.03
9	CH <sub>3</sub> /F	-4.73	-3.45	1.29	-5.15	-2.85	2.30	-4.48	-3.58	0.90
10	F/OCH <sub>3</sub>	-4.74	-3.29	1.46	-5.15	-2.67	2.48	-4.51	-3.46	1.05
11	CH <sub>3</sub> /H (F-T)	-4.85	-3.55	1.30	-5.30	-2.96	2.34	-4.66	-3.67	0.99
12	CH <sub>3</sub> /H (NTZ)	-4.72	-3.53	1.19	-5.12	-2.96	2.16	-4.46	-3.61	0.85
13	H	-4.93	-3.49	1.44	-5.31	-2.86	2.45	-4.43	-3.68	0.75
14	F	-5.19	-3.63	1.56	-5.61	-3.04	2.57	-4.74	-3.88	0.86
15	OCH <sub>3</sub>	-4.75	-3.26	1.49	-5.15	-2.65	2.50	-4.24	-3.37	0.87
16	CN	-5.73	-4.40	1.34	-6.14	-3.83	2.30	-5.57	-4.61	0.96
17	H	-4.87	-2.44	2.42	-5.36	-1.72	3.63	-4.48	-3.12	1.36
18	F	-5.00	-2.64	2.36	-5.53	-1.94	3.59	-4.75	-3.39	1.36
19	OCH <sub>3</sub>	-4.19	-2.22	1.97	-4.74	-1.53	3.21	-3.96	-2.81	1.15
20	CH <sub>3</sub>	-4.67	-2.36	2.31	-5.16	-1.66	3.50	-4.34	-3.01	1.33
21	CN	-5.59	-3.54	2.05	-6.08	-2.89	3.19	-5.29	-4.13	1.16
22	PNTz4T-2OD	-4.88	-3.42	1.46	-5.32	-2.88	2.49	-4.26	-3.49	0.77
23	PfBT4T-2OD	-4.66	-3.38	1.28	-5.03	-2.78	2.25	-4.29	-3.46	0.83

<sup>a</sup> $R_1$  and  $R_2$  are the functional groups in the D and A units, respectively. Structures 11 and 12 have the same functional groups as structure 2, but the first one has been fluorinated on the T rings in position 3 and 4, while in the second one the BTZ moiety has been substituted with NTZ. Note that structures from 13 to 16 (BTZ-T, A units only), and from 17 to 21 (BDT-T, D units only) present one functional group only. Structures 22 and 23 are not part of the PBTZT-*stat*-BDTT-8 series, and therefore, only their names are shown.

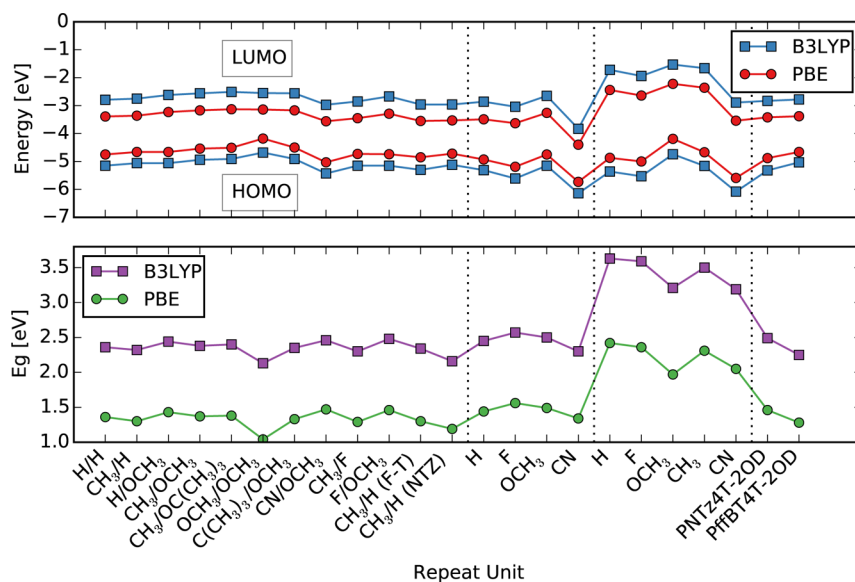


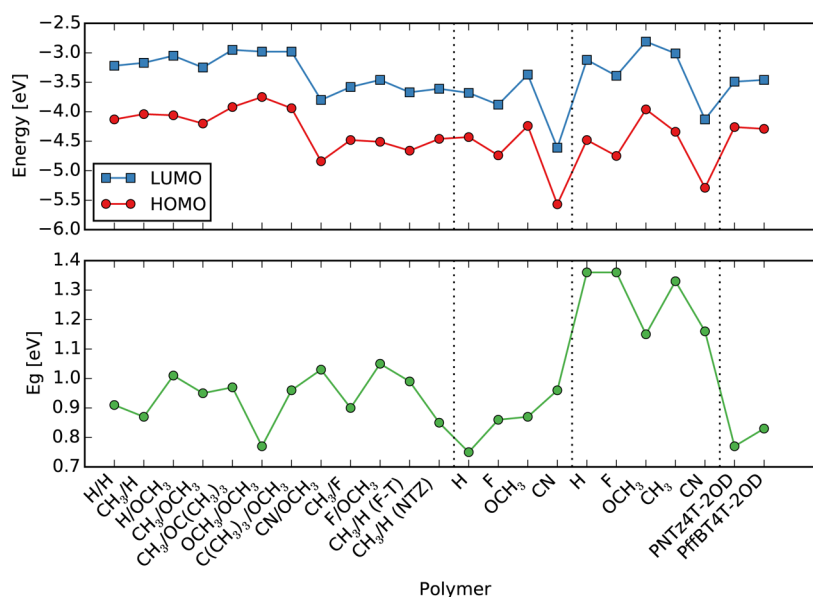
Figure 3. Graphical comparison of HOMO, LUMO and  $E_g$  of repeat units obtained with ONETEP using PBE and B3LYP functionals.

B3LYP functional is to shift HOMO (LUMO) levels toward lower (higher) values, but keeping the overall trend unaltered. In particular the shift in HOMO energies ranges between 0.4 and 0.5 eV, while for LUMO energies the range is between 0.6 and 0.7 eV.

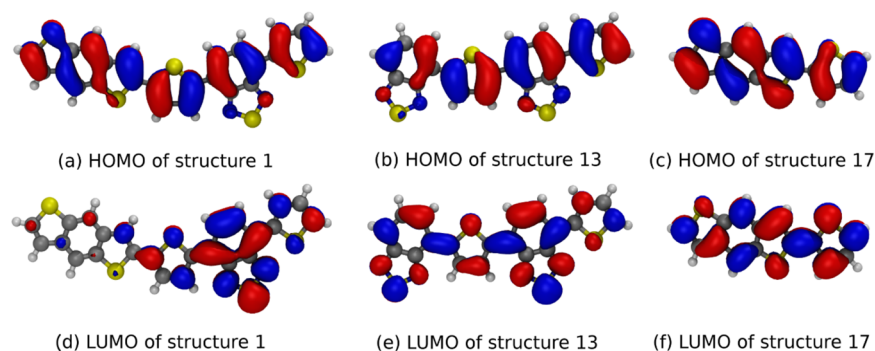
Considering now the polymers, band gaps of structures from 1 to 12 range between 0.77 eV for structure 6 ( $R_1 = R_2 =$

OCH<sub>3</sub>) to 1.05 eV for structure 10 ( $R_1 = F, R_2 = OCH_3$ ). Even though chemistry considerations would suggest that the strategy to reduce the band gap would be to increase the HOMO energy by attaching an electron-donating group on the D unit, and to reduce LUMO energy by attaching an electron-withdrawing group on the A unit, results show that this is not necessarily true: structure 6, for example, presents the lowest





**Figure 4.** Graphical comparison of HOMO, LUMO, and  $E_g$  of polymers obtained with ONETEP using the PBE functional with PBC.



**Figure 5.** Isovalue surface plots of HOMO and LUMO of repeat units 1, 13, and 17.

band gap in the series despite having electron-donating methoxy groups in both D and A units. For structure 9 instead, with methyl groups attached to the D unit and fluorine atoms in the A unit, functionalization does not seem to have any appreciable effect, at least in the band gap, which is essentially the same as structure 1 with both hydrogens as  $R_1$  and  $R_2$ , although both HOMO and LUMO energies are shifted toward lower values. The opposite behavior was observed for HOMO and LUMO levels of structure 6, since both of them increased after functionalization, although the effect was more intense for the HOMO (from  $-4.13$  to  $-3.75$  eV) rather than for the LUMO (from  $-3.22$  to  $-2.98$  eV); this is the reason why structure 6 presents the lowest band gap in the series. Structures 5 and 7, both with bulky substituents, present almost the same band gap as structure 4, despite the HOMO and LUMO energies being different by  $\sim 0.3$  eV: this effect is probably due to the steric hindrance caused by the *t*-butyl and *t*-butoxy groups that leads to slightly nonplanar structures, and therefore a lower degree of conjugation and higher HOMO and LUMO energies. Fluorination on thiophene does seem to have an effect (in contrast to what can be observed for repeat units), causing an increase in the band gap of 0.12 eV and an appreciable decrease in both HOMO and LUMO levels, when comparing structure 2 with 11. The replacement of BTZ with NTZ also has an effect, in particular on the HOMO and LUMO energies: when comparing structures 2 and 12 (which

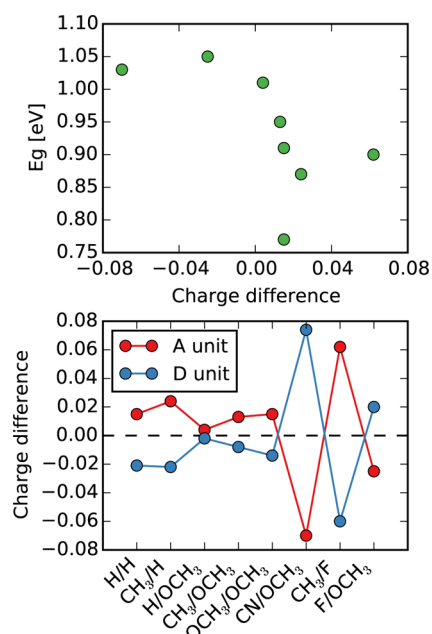
have the same functional groups), it is clear how naphthothiadiazole shifts both the frontier orbitals toward lower values, while the band gap does not seem to be strongly affected. Structure 14, formed by an alternating fluorinated benzothiadiazole and a single thiophene unit, presents almost the same band gap as structure 23 (PffBT4T-2OD), that consists of the same fluorinated benzothiadiazole unit, but alternated with four thiophene units, despite both HOMO and LUMO energies being higher in the latter. Of great importance is also the HOMO energy offset between the donor and the acceptor blocks: as previously mentioned, a well-performing material must have nonvanishing hole mobility, and therefore valence bands of acceptor and donor blocks should be similar. Structure 13 ( $R = H$ ), for example, matches quite well with structures 17 ( $R = H$ ) and 20 ( $R = CH_3$ ), structures 14 and 18, both with  $R = F$ , have almost exactly the same HOMO energy, and structure 15 ( $R = OCH_3$ ) matches reasonably well with structure 20 ( $R = CH_3$ ). Structure 16 ( $R = CN$ ) presents a really deep HOMO level and it does not seem to match with any of the other blocks, perhaps only with structure 21, which has the same CN functional group, although the difference in energy is around 0.30 eV.

The localization of frontier orbitals in both repeat units and polymers was also investigated, as shown in Figure 5: the isosurface plots of HOMO and LUMO for structures 1–12 show a general pattern where HOMO is mostly delocalized

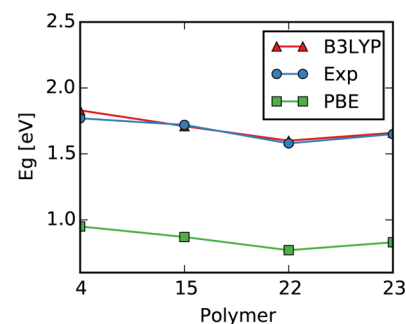
throughout the repeat units (polymers) backbone, while LUMO is mostly localized on the acceptor units (i.e., BTZ and NTZ), and this is typical for most of the D–A polymers present in literature.<sup>31,32</sup> Some functional groups, however, are able to significantly influence the orbitals' localization, and this is the case of structures 6 and 8. Structure 6 presents a localized HOMO on the BDT donor unit due to the electron-donating OCH<sub>3</sub> group attached to it, while in structure 8, that differs from the previous one only for the CN group attached to the D unit, both HOMO and LUMO are delocalized along the chain; this may suggest that for structure 8 an intramolecular charge separation is less likely to be observed, if observed at all. Structures 13–16, composed of alternated benzothiadiazole and thiophene moieties, all show delocalized HOMO and LUMO, regardless of the functional group. This behavior is not typical for D–A copolymers (in this case, thiophene rings can be regarded as donor groups), as for these structures, frontier orbitals are as delocalized as in homopolymers. The same localization can be observed for structures 17–21 composed of benzodithiophene and thiophene.

Also, Mulliken population analysis was performed in order to investigate if and how charge redistribution between D and A blocks may correlate with the band gap trend previously shown and discussed. Charges were computed on a set of polymers consisting of eight different D–A structures and the corresponding D- and A-only structures; the charge difference between A and D blocks when bonded together and when bonded with themselves was computed in order to provide an estimate of how electrons redistribute after copolymerization. The charge redistribution was then defined as the sum of the charge difference of D and A blocks separately (note that the total sum of the charge difference between D and A blocks is zero, as expected). From the bottom plot of Figure 6 it can be observed that in most of the cases electrons move from the donor to the acceptor unit, even when blocks have the same functional groups (as in the H/H and OCH<sub>3</sub>/OCH<sub>3</sub> cases), showing that the BTZ-T moiety has a natural tendency of attracting electrons. However, if a strong electron withdrawing group, like F or CN, is attached to the BDT-T moiety, electrons flow in the opposite direction, that is from the A to the D block. The top plot of Figure 6 shows the band gaps of the eight D–A copolymers as a function of the charge redistribution: it is evident how there does not appear to be any kind of correlation between these two quantities.

Experimental values of polymer band gaps, where available, were also compared with the results both obtained from simulations using PBE and B3LYP functionals. While PBE results were computed under periodic boundary conditions, B3LYP values were obtained with open boundary conditions due to the impossibility of performing periodic calculations with hybrid functionals in the current version of ONETEP, as previously mentioned in Methods; therefore, the B3LYP results refer to oligomers consisting of 5–6 repeat units each. The number of repeat units for this set of oligomers was chosen after performing a convergence analysis of the band gap against the number of units, so that the error in the band gap of these oligomers was between 0.04 and 0.03 eV. Figure 7 and Table 3 show that, despite their absolute values being different due to the use of PBE and B3LYP functionals, the band gaps obtained with ONETEP follow the same trend as in experiments. As expected, the PBE provides an underestimate of the band gap, while the hybrid functional is a slight overestimate, with the only exception of structure 15 whose computed band gap is



**Figure 6.** Electron redistribution as a function of the chemical substituents (bottom graph), and band gap of copolymers as a function of the electron redistribution (top graph). The electron redistribution was defined so that positive values correspond to an increase in the number of electrons, while negative values correspond to a decrease. The names of the polymers are in the D/A format, that is, the first functional group is the one attached to the D unit, while the second is the one attached to the A unit.



**Figure 7.** Comparison between theoretical and experimental band gaps of polymers.

**Table 3. Comparison between Numerical Values (in eV) of Theoretical and Experimental  $E_g$  of Polymers**

polymer	exp	PBE	B3LYP
4	1.77 <sup>14</sup>	0.95	1.83
15	1.72 <sup>33</sup>	0.87	1.71
22	1.58 <sup>a</sup>	0.77	1.60
23	1.65 <sup>b</sup>	0.83	1.66

<sup>a</sup>See ref 21. <sup>b</sup>See ref 20.

0.01 eV lower than the experimental one; however, this deviation is small and within the experimental error. These two functionals provide a lower and a (slight) upper bound, respectively, for the experimental band gaps, even if it is worth remembering that it is not certain that B3LYP will produce overestimates of those quantities: formally Hartree–Fock calculations would provide the upper bound due to the way that the exchange interaction is calculated; B3LYP contains

only a fraction of Hartree–Fock exchange so this effect is moderated. Finally, it is also worth noting that the difference between the experimental and the theoretical values is rather constant, given the experimental error in the determination of optical band gaps.

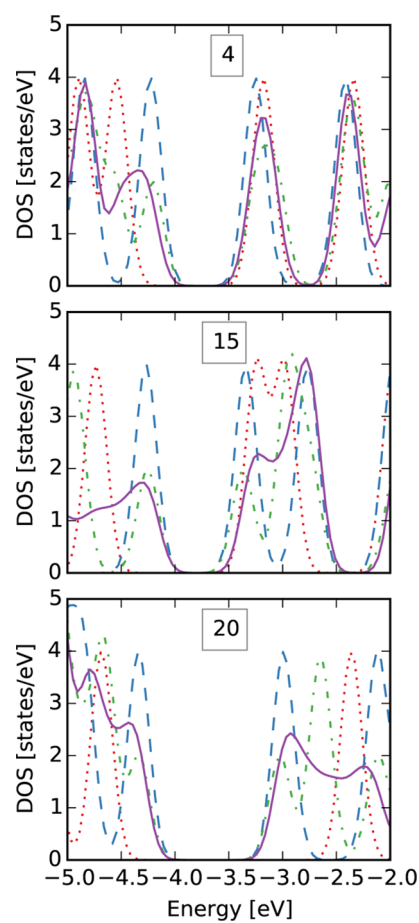
**Long Chain Oligomers.** As previously mentioned, the five long chain oligomers have all the same functional groups in  $R_1$  and  $R_2$  positions ( $\text{CH}_3$  and  $\text{OCH}_3$ , respectively), and therefore, the first three oligomers taken from structures 4 ( $R_1 = \text{CH}_3$ ,  $R_2 = \text{OCH}_3$ ), 15 ( $R = \text{OCH}_3$ ), and 20 ( $R = \text{CH}_3$ ) were built, and the results obtained after geometry optimization were compared with the corresponding ones obtained in periodic boundary conditions, as shown in Table 4. The oligomers of

**Table 4. Energies of Oligomers Obtained after Geometry Optimization, Compared with the Ones Obtained with Periodic Boundary Conditions**

	HOMO (eV)	LUMO (eV)	$E_g$ (eV)
		structure 4	
PBC (1 unit)	−4.20	−3.25	0.95
PBC (2 units)	−4.20	−3.26	0.94
oligomer	−4.20	−3.28	0.92
		structure 15	
PBC (1 unit)	−4.28	−3.34	0.94
PBC (2 units)	−4.24	−3.38	0.87
oligomer	−4.20	−3.34	0.86
		structure 20	
PBC (1 unit)	−4.34	−3.01	1.33
PBC (2 units)	−4.33	−3.02	1.31
oligomer	−4.32	−3.03	1.29

structures 4, 15, and 20 are composed of 19, 18, and 20 repeating units, respectively, and Figure 8 shows a comparison between structure 5 oligomer, a dimer in periodic boundary conditions, and finally a single repeat unit. The HOMO and LUMO energies of the oligomers were found to be in very good agreement with the ones obtained in PBC; this is further confirmation that these polymers have very flat bands: for structures 1–11 and 17–21, only one repeating unit in the unit cell is enough to obtain reasonably converged energies, while for structures 13–16, at least two units are necessary.

Figure 9 shows a comparison between density of states (DOS) centered around the energy gaps of single repeat units, polymers obtained in PBC using just one repeating unit, polymers in PBC but with two repeating units, and finally oligomers with 19, 18, and 20 units. For each of the structures it was observed that the DOS between repeat units and oligomers



**Figure 9.** Comparison between density of states of repeat units (dotted red line), polymers obtained in PBC with 1 single unit in the repeating cell (dashed blue line), polymers obtained in PBC with 2 units (dotted and dashed green line), and oligomers (solid purple line) of structures 4 (top), 15 (middle), and 20 (bottom), respectively.

are significantly different: not only the HOMO and LUMO energies get closer because of the effect of conjugation, leading to a narrowing in the energy gaps, but a broadening of the states can also be observed, especially for structures 15 and 20. This is consistent with the fact that while for single repeat units the energy levels are well-defined and separated, for oligomers they become significantly closer to one another, and eventually a semiconductor band structure is observed. When comparing DOS of oligomers and polymers obtained in PBC it is also clear that, in general, a repeating cell with just one unit (or two units,



**Figure 8.** Ball and stick representation of structure 4 oligomer, dimer in PBC, and repeat unit.

in the case of structure 15) is enough to obtain converged numerical values of the HOMO and LUMO energies, even if the density of states overall are clearly different.

Results of the ground state energies of oligomers obtained with different reactivity ratios are shown in Table 5. Structures

**Table 5. Energies of Oligomers Obtained after Geometry Optimization with Different Reactivity Ratios for D and A Blocks<sup>a</sup>**

	no. atoms	HOMO (eV)	LUMO (eV)	$E_g$ (eV)
pure D	1046	-4.32	-3.03	1.29
pure A	1042	-4.20	-3.34	0.86
alternating D–A	1047	-4.20	-3.28	0.92
D10-A0.1	1156	-4.29	-3.29	1.00
D5-A0.2	1150	-4.28	-3.26	1.02
D0.2-A5	1069	-4.23	-3.36	0.87
D0.1-A10	1054	-4.22	-3.36	0.87
D1-A1	1096	-4.25	-3.34	0.91

<sup>a</sup>Pure D and pure A refer to BDT-T oligomer and BTZ-T oligomer, respectively.

D10-A0.1 and D5-A0.2, composed mostly of D blocks, present HOMO energies close to the one of the pure D oligomer, while LUMO energies are closer to the one of the pure A structure; the band gaps, with values of 1.00 and 1.02 eV, respectively, present intermediate values between the ones of pure D and A structures, and they are both wider than the one of the alternating D–A oligomer. Structures D0.1-A10 and D0.2-A5, that consist mostly of A blocks, on the other hand, present both HOMO and LUMO energies that are closer to the one obtained for the BTZ-T oligomer; both energy gaps in turn present almost the same values as the one of the BTZ-T oligomer and, therefore, smaller in magnitude than the one of the alternated D–A oligomer. Finally, structure D1-A1 (i.e., a totally random copolymer) presents a HOMO level that is almost the average between that predicted for the pure D and A oligomers, while the LUMO energy is exactly the same as the one of the pure A structure. Small changes were observed in the HOMO and LUMO energies, in the order of <0.05 eV, when comparing the values between the alternated copolymer and the totally random one; the same observation was found to be true also for the energy gap, with a change of ~0.01 eV. It seems, therefore, that the “sequence effect”<sup>34</sup> (i.e., the order of the blocks in the chain) does not play an important role, at least for this class of materials.

Frontier orbitals of all the oligomers studied present the same localization behavior as D–A polymers: HOMO is mostly delocalized on the oligomer backbone, while LUMO is strictly localized on the acceptor groups of the chain.

## CONCLUSIONS

D–A copolymers are widely used in the field of organic photovoltaics because of their versatility: HOMO and LUMO energies, as well as the band gap, can be carefully tuned in different ways in order to obtain materials with suitable and desired optical properties (such as a low band gap), with the aim of boosting the efficiency of this kind of devices. This study in particular focused on the effect of composition on a class of statistical copolymers, and the model structures used in this paper consisted of analogues and blocks of the high-performing PBTZT-*stat*-BDTT-8 developed by Merck.

The effect of composition was studied by modifying the D/A block ratio in the polymer chain, without changing the functional groups. The block ratio was modified by setting different values of reactivity ratios, and the final compositions of the oligomers were obtained by using the Mayo–Lewis equation as implemented in the Materials Studio software. This study was enabled by the linear-scaling reformulation of DFT in the ONETEP code, that allowed calculation at a far larger scale than possible before: it was then possible to study a series of long chain oligomers composed of more than a thousand atoms, allowing us to obtain well converged energy levels, which would have been impossible otherwise (previous theoretical works on this subject made use only of small tetramers<sup>11</sup>). Our results show that the composition plays a very important role in the determination of the electronic properties of a D–A copolymer: when varying the D/A block ratio, changes in the energy gap of the same material were observed and were in the order of ~0.1 eV, an effect that is to some extent comparable to changing the functional groups. The band gap of the studied material ranged between ~1 eV (in the case where more D units were present in the chain) and 0.87 eV (in the opposite case, where a higher number of A units was present in the chain). From these results it is clear that modifying the block ratio of the polymer (in other words, modifying the synthesis strategy) leads to structures with significantly different electronic properties, without the need to change the functional groups. In addition, at least for this class of statistical copolymers, the sequence effect does not seem to significantly affect the HOMO, LUMO, and band gap, since only a small change in these values was observed when comparing the energies of an alternating D–A oligomer with a completely random one, built by assigning the same reactivity ratios for both blocks.

An extensive study on the electronic properties of this class of materials was also performed: in order to understand how the electronic structure of PBTZT-*stat*-BDTT-8 is susceptible to functionalization, the effect of a number of substituents attached to its donor and acceptor units was investigated. Different functional groups were tested, both electron-withdrawing and electron-donating. In general, electron-withdrawing groups were found to cause a decrease in both HOMO and LUMO energies and in both A and D units, while electron-donating groups on the contrary showed the opposite effect. However, when combining the donor and the acceptor units together, it was hard to rationalize the effect of specific substituents in terms of HOMO and LUMO energies: this suggests that it is unlikely to know a priori using chemical intuition how a functional group can influence the energy levels of a D–A copolymer, and therefore, computational modeling is a useful and essential tool. This finding is also supported by the fact that no correlation could be found between the band gaps and the charge redistribution of the materials studied. The results were also supported by comparing computed energy gaps, using both hybrid and GGA functionals, with the corresponding experimental ones present in literature, and despite the obtained theoretical values with PBE being smaller in magnitude than the experimental ones (or slightly higher and/or comparable in the case of B3LYP), we were able to accurately reproduce the experimental trend.

The results shown in this paper can be used as a starting point for the design of efficient D–A statistical copolymers with suitable electronic properties for OPV devices. However, more in-depth studies need to be done in order to address more



complex problems regarding these kinds of materials, such as the effect of chain–chain stacking and the interaction between these polymers and fullerenes.

## ■ ASSOCIATED CONTENT

### 5 Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jpcc.6b10851.

Extrapolation of band gaps by linear and quadratic fit for a set of structures (4, 15, 22, 23), as a comparative example to results obtained with PBC and, in some cases, to results of the long chain oligomers (PDF).

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### Notes

The authors declare no competing financial interest.

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