

PCCP

Accepted Manuscript



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

Accepted Manuscripts are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this *Accepted Manuscript* with the edited and formatted *Advance Article* as soon as it is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.

Threshold-like Complexation of Conjugated Polymers with Small Molecule Acceptors in Solution within the Neighbor-Effect Model

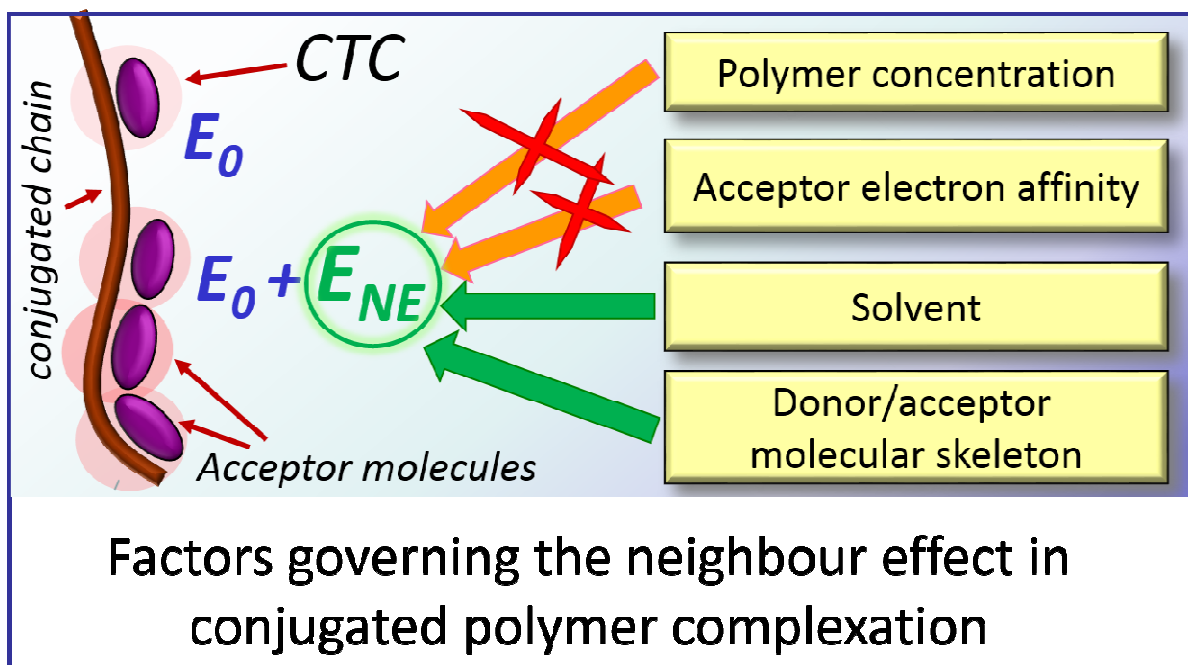
Andrey Yu. Sosorev¹, Olga D. Parashchuk¹, Sergey A. Zapunidi¹, Grigoriy S. Kashtanov¹, Ilya V. Golovnin¹, Srikanth Kommanaboyina², Igor F. Perepichka², and Dmitry Yu. Paraschuk^{1*}.

¹ Faculty of Physics and International Laser Center, M.V. Lomonosov Moscow State University, Moscow 119991, Russia.

² School of Chemistry, Bangor University, Bangor LL57 2UW, UK

*E-mail paraschuk@gmail.com

TOC figure



Abstract

In some donor–acceptor blends based on conjugated polymers, a pronounced charge-transfer complex (CTC) forms in the electronic ground state. In contrast to small-molecule donor-acceptor blends, the CTC concentration in polymer:acceptor solution can increase with acceptor content in a threshold-like way. This threshold-like behavior was earlier attributed to the neighbor effect (NE) in the polymer complexation, i.e., next CTCs are preferentially formed near the existing ones;

however, the NE origin is unknown. To address the factors affecting the NE, we record the optical absorption data for blends of the most studied conjugated polymers, poly(2-methoxy-5-(2-ethylhexyloxy)-1,4-phenylenevinylene) (MEH-PPV) and poly(3-hexylthiophene) (P3HT), with electron acceptors of fluorene series, 1,8-dinitro-9,10-antraquinone (**DNAQ**), and 7,7,8,8-tetracyanoquinodimethane (**TCNQ**) in different solvents, and then analyze the data within the NE model. We have found that the NE depends on the polymer and acceptor molecular skeletons and solvent, while it does not depend on the acceptor electron affinity and polymer concentration. We conclude that the NE operates within a single macromolecule and stems from planarization of the polymer chain involved in the CTC with an acceptor molecule; as a result, the probability of further complexation with the next acceptor molecules at the adjacent repeat units increases. The steric and electronic microscopic mechanisms of NE are discussed.

Introduction

Blends of conjugated polymers (CPs) with small-molecule acceptors are the heart of the state-of-the-art plastic solar cells. In some of the blends, a pronounced charge-transfer complex (CTC) is formed between the polymer donor and the acceptor molecules in the electronic ground state.¹⁻⁴ The CP CTCs can significantly absorb in the red/near-IR regions^{1, 5, 6} and generate mobile charges^{7, 8} making them promising for photovoltaics. As the lowest electronic excited states in the donor-acceptor blends, the excited CTCs (usually called charge-transfer states) are key intermediates on the pathway from excitons to free charges⁹ even in CP blends with fullerenes where the CTC is very weak.¹⁰⁻¹² As a result, the CTCs strongly affect the solar cell performance.¹³⁻¹⁵ The ground-state donor-acceptor interaction in CTCs can also influence the conformation of the polymer chains,^{6, 16} the phase separation in the blend,^{3, 17} and enhance the photooxidation stability.^{8, 18} All these CTC features could be exploited to improve the performance of organic solar cells.⁴

CTCs in small-molecule donor-acceptor blends have been thoroughly investigated. However, if one of the CTC constituents, e.g. donor, is a CP, its macromolecular nature and electron delocalization over the conjugated segment can significantly affect the CTC formation and properties. Specifically, the CTC concentration in solution of poly(2-methoxy-5-(2-ethylhexyloxy)-1,4-phenylenevinylene), MEH-PPV, and 2,4,7-trinitrofluorenone, **TNFon**, increases in a clear threshold-like fashion:⁶ when the acceptor concentration is below a threshold value, the CTC concentration is low; however, when the threshold is exceeded, an intensive CTC formation starts. This is contrasting to the small-molecule blends, where the CTC concentration increases gradually with an acceptor addition, and the plot of the CTC concentration versus the acceptor concentration is convex. To explain the threshold-like CTC formation in MEH-PPV:**TNFon** blends, the neighbor effect (NE) model was recently proposed.^{6, 19} According to this model, the CTC binding energy increases if two or more acceptor molecules are complexed with the polymer chain nearby each other. **Fig. 1** illustrates this NE effect: the binding energy of an isolated (individual) CTC is E_0 , and it increases by E_{NE} or $2E_{NE}$ ($E_0, E_{NE} > 0$) if one or both of the neighboring segments are also complexed, respectively. Hence, successive CTCs are preferentially formed near the existing ones, i.e. the CTC formation is a cooperative phenomenon. For MEH-PPV:**TNFon** blends, it was found that E_{NE} can be comparable to E_0 so that the NE can play a key role in the CP complexation.¹⁹ The NE should result in formation of intrachain CTC aggregates that can explain photoluminescence quenching in MEH-PPV:**TNFon** blends.²⁰ Furthermore, these aggregates seem to be inherited in film allowing explanation of ordered polymer nanodomains in MEH-PPV:**TNFon** blends.²¹ Therefore, the NE can lead to blend self-organization. Unraveling the mechanisms of cooperative phenomena and self-organization in polymers is a long-standing challenge in the polymer science; moreover, the cooperative effects in CPs involving their π -conjugated system are practically not studied. Accordingly, the NE in CP complexation is worth detailed investigation. Specifically, it is not known whether the NE is a general feature of CTC formation in CP donor-acceptor blends, or it

occurs only in MEH-PPV:TNFon blend. Consequently, the physical factors that drive the NE, and its microscopic mechanism should be addressed.

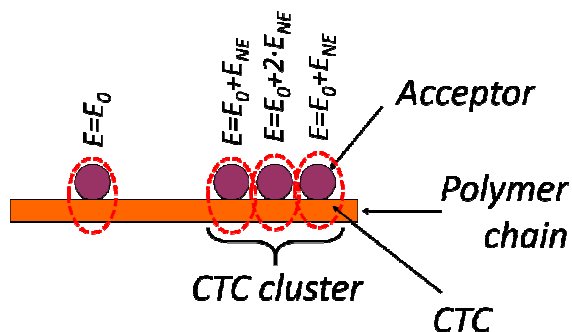


Fig. 1. Schematics of the neighbor effect.

There are several factors that can influence the CTC formation in solution. According to the Mulliken model,²² the CTC binding energy depends on the difference between the donor ionization potential corresponding, in one-electron model, to the highest occupied molecular orbital (HOMO) energy, and the acceptor electron affinity corresponding to the lowest unoccupied molecular orbital (LUMO) energy. This energy difference is called the “effective HOMO–LUMO gap”.²³ The lower the effective HOMO(D) – LUMO(A) gap, the higher the CTC binding energy. As a result, for the same donor and various acceptors, the binding energy should depend on the acceptor electron affinity. The second main factor that determines the CTC binding energy in the Mulliken model is the donor-acceptor electronic coupling, H_{DA} , which in its turn depends on the compatibility of the donor and acceptor π -conjugated systems at their molecular skeletons. In addition, since the CTC formation in solution is affected by the donor-solvent and acceptor-solvent interactions, the solvent is also expected to affect the CTC binding energy. Finally, as shown in Ref. 24, the CTCs serve as interchain links, and it is not clear whether CTCs can be formed on a single chain. Therefore, the CTC formation could depend on the polymer concentration.

In this work, we address the influence of the effective HOMO–LUMO gap, solvent, polymer concentration, and molecular skeletons of the donor and acceptor on the CTC formation in blends of CPs with small-molecule acceptors. We study CTC optical absorption in blended solutions of MEH-PPV and poly(3-hexylthiophene) (P3HT) with various electron acceptors: acceptors of the fluorene series, 1,5-dinitro-9,10-anthraquinone (**DNAQ**), and 7,7,8,8-tetracyanoquinodimethane (**TCNQ**). Then we fit the data by the analytical NE model for CP complexation¹⁹ and obtain the isolated CTC binding energy, E_0 , and the NE energy gain, E_{NE} , as fitting parameters. Finally, we investigate how E_0 and E_{NE} are affected by the factors mentioned above and discuss the possible microscopic mechanisms of the NE in CP complexation.

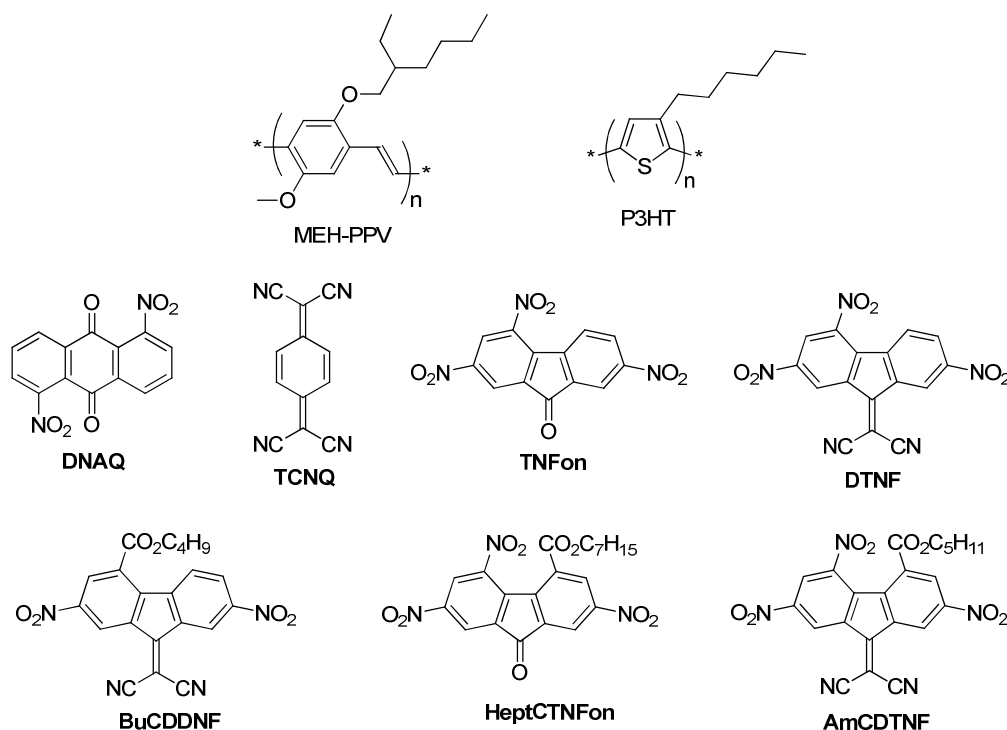


Fig. 2. Structural formulas of studied materials.

Experimental

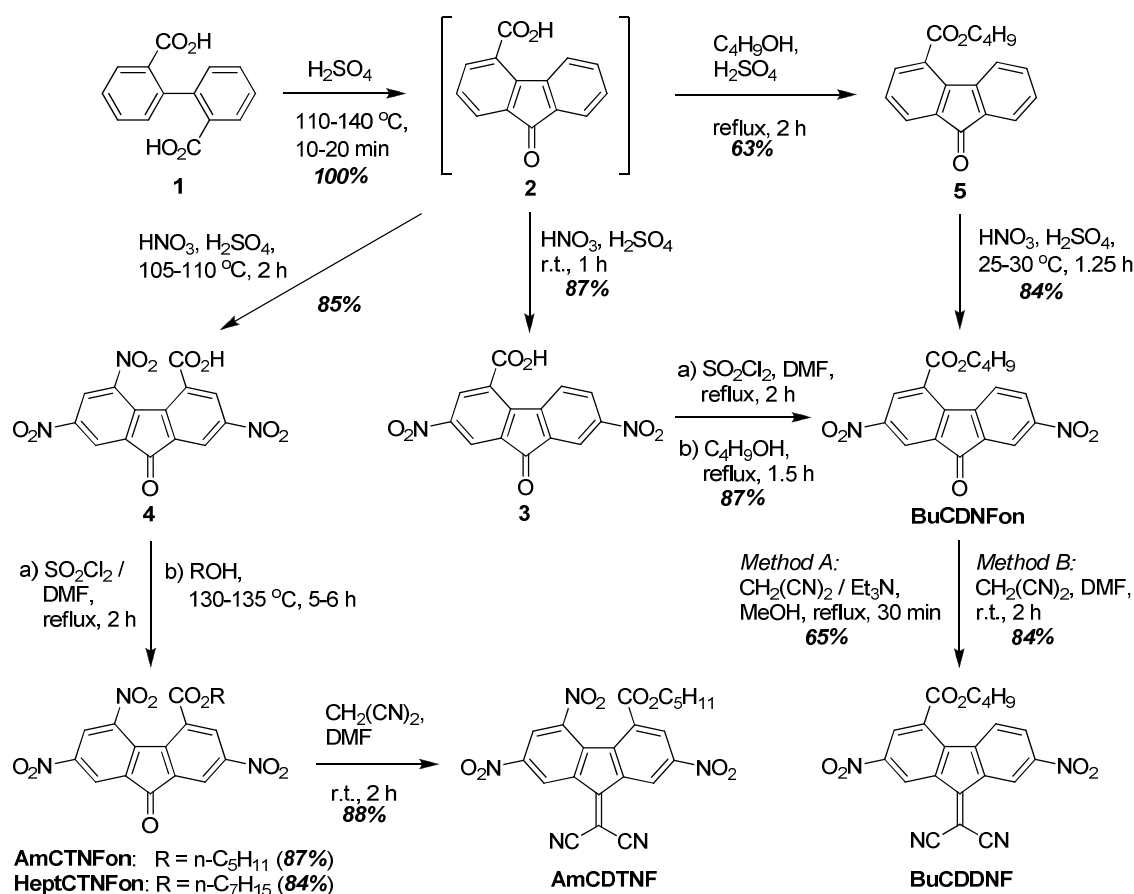
Materials. Fig. 2 shows the structural formulas of studied polymer donors and molecular electron acceptors used in this study. MEH-PPV ($M_w = 86,000$, $M_n = 420,000$, Sigma-Aldrich) and P3HT (regioregular electronic grade, #4002-EE Reike Metals) were used as received. TCNQ was recrystallized twice from acetonitrile. Various fluorene-type electron acceptors have been synthesized and used in this study. These acceptors contain from two to three nitrogroups (NO_2 in positions 2,4,7-) in their structure and carbonyl oxygen atom or dicyanomethylene group at the C-9 position. In addition, some acceptors contained $-\text{CO}_2\text{R}$ ester group with long chain solubilizing linear alkyl substituents $\text{R} = n\text{-C}_4\text{H}_9$, $n\text{-C}_5\text{H}_{11}$, $n\text{-C}_7\text{H}_{15}$ (see Supporting Information for the details of their synthesis). These variations in the chemical structure of the acceptors allowed to vary their electron affinities and solubility. 2,4,7-Trinitrofluorenone (TNFon) and 2,4,7-trinitro-9-dicyanomethylene-fluorene (DTNF) were synthesized according to procedures described previously.²⁵

Computational procedures. DFT computations of the geometries of studied electron acceptors were carried out with the Gaussian 09²⁶ package of programs by using Pople's 6-31G split valence basis set supplemented by d-polarization functions and diffusion functions for heavy atoms. Becke's three-parameter hybrid exchange functional^{27, 28} with the Lee–Yang–Parr gradient-corrected correlation functional (B3LYP)²⁹ were employed. The restricted Hartree-Fock formalism was used. No constraints were used and all structures were free to optimize in a chlorobenzene or toluene solution using the polarizable continuum model (PCM).³⁰ Thus, the geometries were optimized at the B3LYP/6-31+G(d) and the electronic structures were calculated at the same level of theory. The electron affinities (EA) were approximated as the negative of the LUMO energies ($EA = -E_{\text{LUMO}}$) in accordance with DFT-Koopman's theorem.

Optical Absorption Spectroscopy. The donor and acceptor were dissolved separately in solvent [chlorobenzene (CB), o-dichlorobenzene (DCB), toluene (Tol), chloroform (CF) or tetrahydrofuran (THF)] and then mixed. The color of the mixture stabilized within ca. 10–20 seconds, the mixture was kept for at least 3–5 min before measurements. The MEH-PPV concentration in CB was kept constant at 1.00 g/l (3.84 mM) except for blends with **TNFon** where it was either 2.00 g/l (7.68 mM) or 0.010 g/l (38.4 μ M). The P3HT concentration was 5.00 g/l (30.1 mM). In solvents effect measurements, the MEH-PPV concentration was kept 2.00 g/l (7.68 mM) in CB or 0.50 g/l (1.92 mM) in other solvents. Electron absorption spectra of mixed solutions were recorded using a fiber-coupled spectrophotometer (Avantes) in quartz cuvettes of 0.10 mm path length (for blends with **TNFon**, **HeptCTNFon**, **BuCDDNF**, **AmCDTNF** and **DNAQ**) or 10 mm path length (for blends with **DTNF** and **TCNQ**, and P3HT:**TNFon**). The CTC absorption coefficients, α_{CTC} = Absorbance/path length [cm], were calculated at 630 nm (for MEH-PPV blends with **TNFon**, **HeptCTNFon**, **BuCDDNF**, **AmCDTNF** and **DNAQ**) or at 800 nm (for MEH-PPV blends with **DTNF** and **TCNQ**, and P3HT:**TNFon** blends), measurements at other wavelengths are given in Supporting Information (SI).

Results

Synthesis of Fluorene Acceptors. Syntheses of fluorene acceptors – derivatives of nitrofluorene-4-carboxylic acids – are depicted in Scheme 1. Diphenic acid (**1**) is quantitatively cyclised by concentrated sulfuric acid into fluorenone-4-carboxylic acid (**2**) at ca. 110–140 °C.^{31, 32} Nitration of the latter by a mixture of fuming nitric and concentrated sulfuric acid results in 2,7-dinitro- (**3**)^{33, 34} or 2,5,7-trinitro-9-fluorenone-4-carboxylic acids (**4**).^{33, 35} From this point of view, an isolation of acid **2** is not necessary and we have elaborated high-yields (>70% after purification) one-pot syntheses of acids **3** and **4** directly from compound **1** by direct adding fuming acid to the solution of **2** formed in sulfuric acid (deep-red solution of protonated form of **2** at its carbonyl oxygen) and performing nitration reactions at room temperature (for **3**) or at 105–110 °C (for **4**) (see Supporting Information). Obtained thus acids **3** and **4** were converted into corresponding esters (**3** \rightarrow **BuCDNFon**; **4** \rightarrow **AmCTNFon**, **HeptCTNFon**) by reaction with thionyl chloride in presence of catalytic amount of *N,N*-dimethylformamide (DMF) following by esterification of formed acyl chlorides with corresponding alcohols (*n*-butanol, *n*-pentanol or *n*-heptanol, accordingly). Ester **BuCDNFon** was also obtained by an alternative route: fluorenone-4-carboxylic acid (**2**) obtained by heating of diphenic acids (**1**) in sulfuric acid was involved *in situ*, without isolation, into the next step of esterification by *n*-butanol to afford *n*-butyl ester **5**. The nitration of the later by a mixture of fuming nitric acid and concentrated sulfuric acid at room temperature gave ester **BuCDNFon** in an excellent yield of 84%, without hydrolysis of the ester group. Conversion of fluorenone **BuCDNFon** into stronger electron acceptor **BuCDDNF** was done by triethylamine-catalyzed condensation with malononitrile in methanol solution, the method which was initially proposed for condensation of polynitro-9-dicyanomethylene fluorenes.^{36, 37} We also applied for this reaction another, more convenient method of condensation, previously proposed by us, which is based on using DMF as a solvent.^{25, 38-41} This method does not require basic catalyst that allows performing the reaction in milder conditions, and it normally gives higher yields and better purity of 9-dicyanomethylene fluorenes. The later method was also used in synthesis of **AmCDTNF** from **AmCTNFon**, and in both cases it gave excellent yields of the products (84–88% after purification).



Scheme 1. Synthesis of fluorene electron acceptors.

DFT Calculations of Orbital Energies and Electron Affinities of Acceptors. DFT computations have been performed at B3LYP/6-31+G(d) level of theory for both geometry optimizations and calculations of electronic structures of all acceptors. The effect of the solvent was incorporated using the polarizable continuum model (PCM) to match the results with spectroscopic measurements in these solvents. Kohn-Sham frontier orbital energies of acceptors (E_{HOMO} and E_{LUMO}) together with HOMO–LUMO energy gaps ($E_g = E_{\text{LUMO}} - E_{\text{HOMO}}$) and electron affinities ($EA = -E_{\text{LUMO}}$, according to DFT-Koopman's theorem) are collated in Table 1 (more details on B3LYP/6-31+G(d) calculations are given in Supporting Information). An incorporation of the solvent effect is important for an accurate estimation of acceptors frontier energy levels and electron affinities, as solvation (from a gas phase to highly polar acetonitrile) changes e.g. LUMO energies of acceptors by 0.2–0.5 eV while the actual solvent effect on EA depends on the structure of the acceptor (Figs. S1 and S2 in SI). In this work we used B3LYP/6-31+G(d) calculated EA values for studied electron acceptors, which was recently shown correlate well ($r = 0.97\text{--}0.98$) with experimental data on EA estimations from cyclic voltammetry experiments in the same solvent (see Fig. S3 in SI).²⁵

Table 1. HOMO and LUMO energy levels, and electron affinities of studied acceptors from DFT calculations.^a

Compound	E_{HOMO} (eV)	E_{LUMO} (eV)	E_g^b (eV)	EA^c (eV)
DNAQ	-8.00	-3.82	4.18	3.82
TCNQ	-7.30	-4.76	2.54	4.76
TNFon	-7.90	-4.12	3.78	4.12
HeptCTNFon	-7.98	-4.21	3.77	4.21
BuCDDNF	-7.71	-4.21	3.50	4.21
DTNF	-7.94	-4.42	3.52	4.42
AmCDTNF	-8.00	-4.49	3.51	4.49

^a DFT B3LYP/6-31+G(d) calculations in chlorobenzene solution using PCM model. Calculations for other solvents (and for the gas phase) are given in SI, Fig. S1.

^b $E_g = E_{\text{HOMO}} - E_{\text{LUMO}}$. ^c $EA = -E_{\text{LUMO}}$.

Charge-Transfer Complexation of MEH-PPV and P3HT Polymers with Molecular Electron Acceptors. Fig. 3 shows absorption spectra of MEH-PPV:**AmCDTNF** (a representative fluorene acceptor), MEH-PPV:**DNAQ**, MEH-PPV:**TCNQ**, and P3HT:**TNFon** blends in chlorobenzene. The spectra of MEH-PPV blends with different fluorene-type acceptors are similar to that of MEH-PPV:**AmCDTNF** and are shown in SI. Upon acceptor addition, new absorption features in the polymer band gap arise in all the blends, and their intensity is increased with the acceptor content. For MEH-PPV blends with **TNFon** and **DNAQ**, this subgap absorption was previously attributed to the CTC.^{1, 4} The MEH-PPV:**TCNQ** blend shows intragap absorption very similar to that observed in Ref. 42 for a slightly different PPV derivative, MDMO-PPV, and this absorption was attributed to CTC. Note that the absorption spectra for different MEH-PPV blends in Fig. 3 show different shapes indicating that the subgap absorption can hardly be assigned to polymer aggregation (for other acceptors see Fig. S4 in SI). In Ref. 42, the subgap absorption in P3HT blends with various acceptors was also attributed to the CTC. We therefore assign the subgap absorption features observed in all the studied blends to the CTC between polymer donors and molecular electron acceptors.

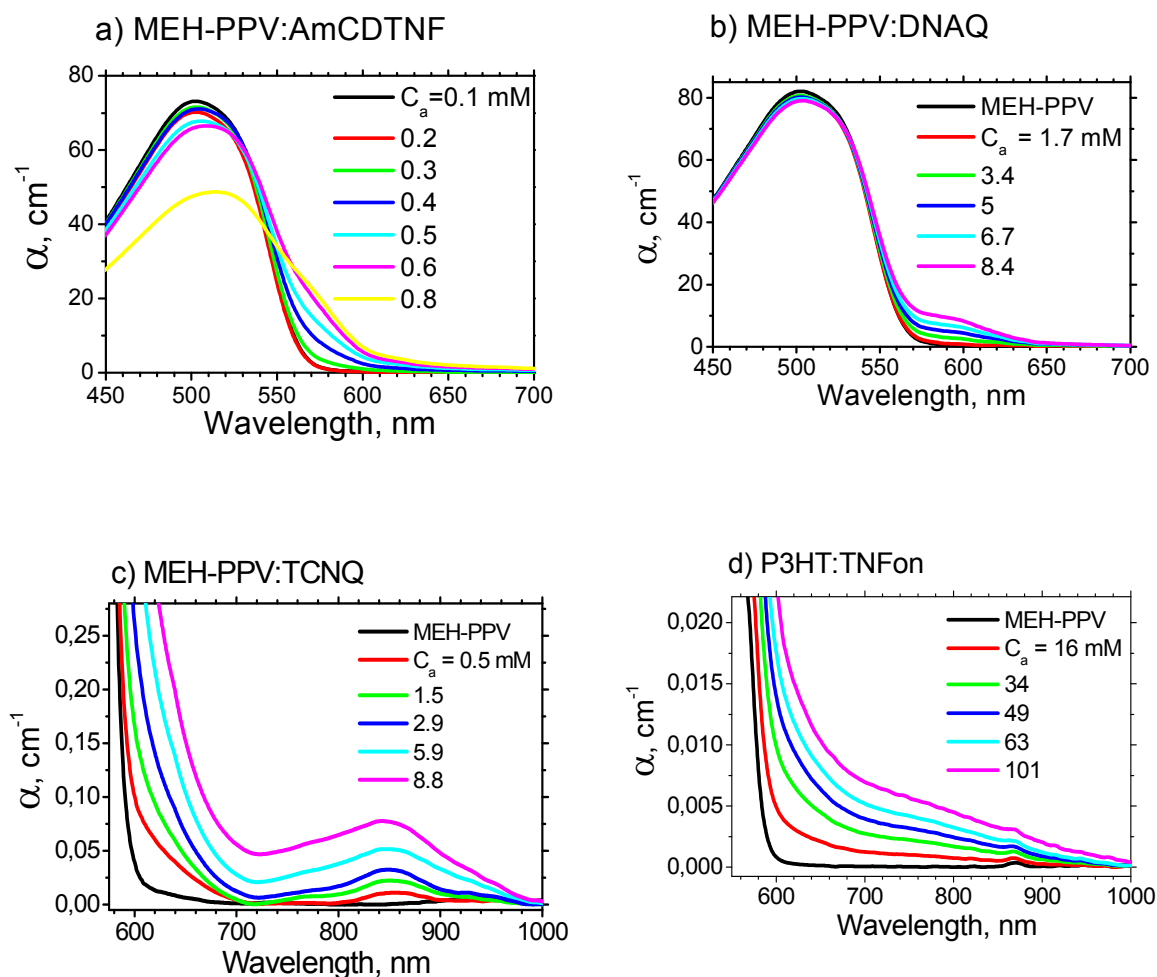
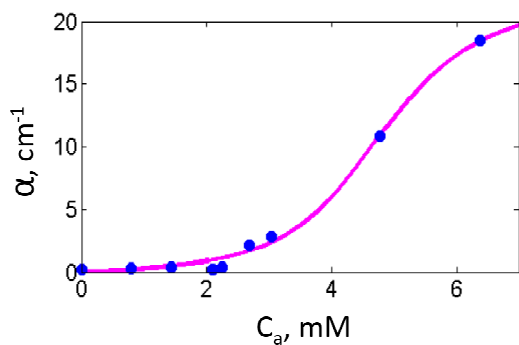


Fig. 3. Absorption spectra of MEH-PPV:**AmCDTNF** (a), MEH-PPV:**DNAQ** (b), MEH-PPV:**TCNQ** (c), and P3HT:**TNFon** (d) blends in chlorobenzene for various acceptor concentrations. A feature at ≈ 870 nm in panel (d) is an instrument artefact. The cuvette path length was 0.10 mm (a, b) or 10 mm (c, d). The MEH-PPV concentration was kept constant at 3.84 mM.

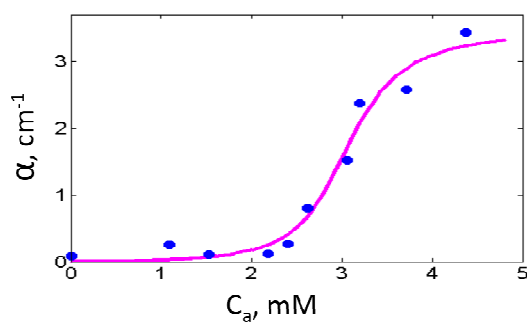
Fig. 4 presents dependencies of the blend absorption coefficients, α , on the acceptor concentration, C_a , for various MEH-PPV:acceptor (a-g) and P3HT:**TNFon** (h) blends at selected wavelengths within the polymer bandgap (the data at other wavelengths are given in Fig. S5, SI). These absorptions correspond to the polymer-acceptor CTC. All the blends, excluding MEH-PPV:**TCNQ** and P3HT:**TNFon**, show similar S-type shapes of $\alpha(C_a)$ dependencies with an inflexion point indicating more or less threshold-type character of polymer complexation. Therefore, we conclude that the threshold-like complexation is not a unique feature of MEH-PPV:**TNFon** blends. The thresholds in the $\alpha(C_a)$ dependence could originate from the threshold-like behavior of the CTC concentration, molar extinction, and/or stoichiometry. For MEH-PPV:**TNFon** blends, the CTC extinction and stoichiometry were shown to be independent of the acceptor content, and the threshold in the CTC absorption was attributed to the threshold-like concentration behavior.⁶ Consequently, we assume that the S-type $\alpha(C_a)$ dependencies in the other MEH-

PPV:acceptor blends are also determined mainly by threshold-like behavior of the CTC concentration, C_{CTC} .

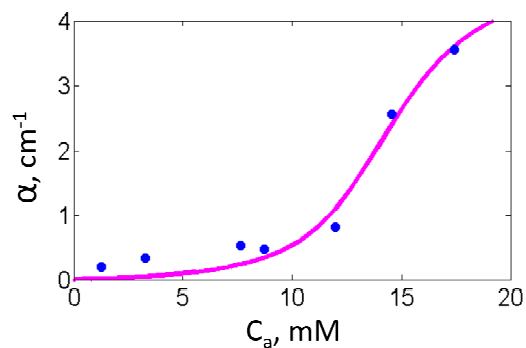
a) MEH-PPV:TNFon



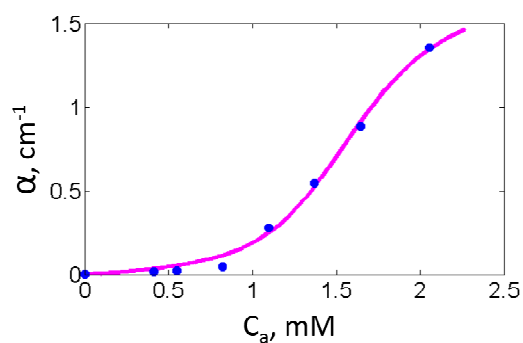
b) MEH-PPV:HeptCTNFon



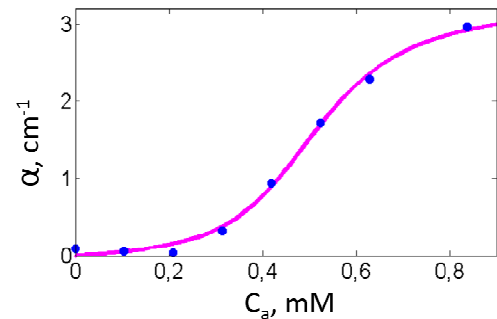
c) MEH-PPV:BuCDDNF



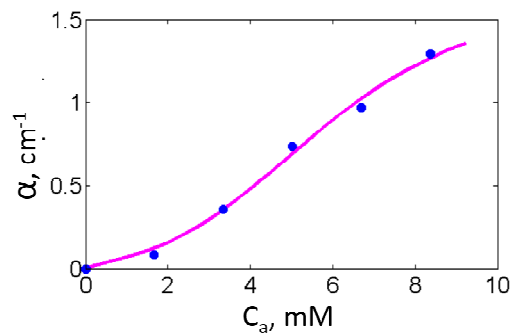
d) MEH-PPV:DTNF



e) MEH-PPV:AmCDTNF



f) MEH-PPV:DNAQ



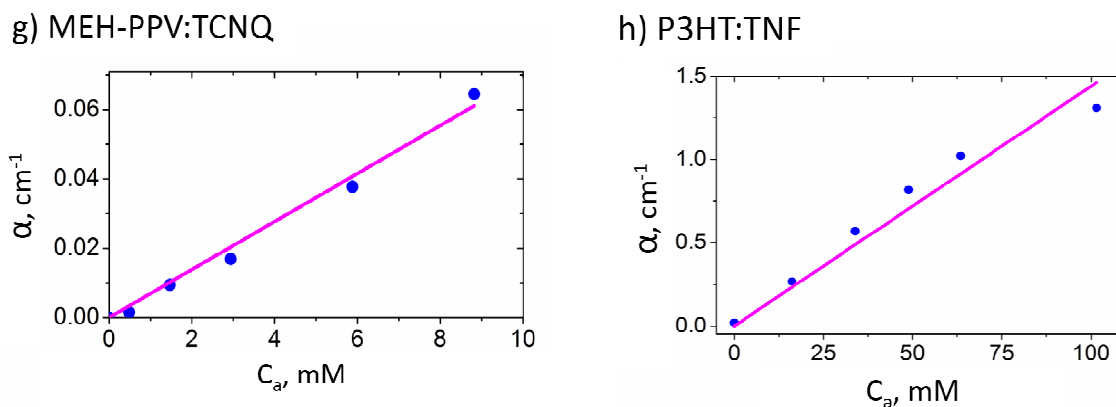


Fig. 4. CTC absorption coefficients (α) as functions of the acceptor concentrations (C_a) for various polymer:acceptor blends in chlorobenzene. The experimental data [absorption at 630 nm (a–c, e, f) or at 800 nm (d, g, h)] are shown by dots. Lines show NE model fits (a–f) and straight-line fits (g,h). The cuvette path length was either 0.10 mm (a–c, e,f) or 10 mm (d, g, h).

While an extension of the C_a range to higher concentrations to reach the saturation of the $\alpha(C_a)$ dependencies in Fig. 4 would be desirable for better fitting the experimental data by the NE model (see below), we are limited by solubility issues. Some acceptors, e.g. DNAQ and TCNQ, have limited solubility in the used solvents. Moreover, at high MEH-PPV and acceptor concentrations, precipitation of the complexed polymer was sometimes observed. This potentially could contribute to the S-shape $\alpha(C_a)$ dependencies. To exclude this possibility, we have performed experiments at a very low MEH-PPV concentration of 0.01 g/L (38.4 μM) and have nicely observed full saturation of $\alpha(C_a)$ dependencies at high acceptor concentrations (see Fig. S7 in SI and discussion of the complexation model below).

Below we discuss how different factors affect the CTC formation in CP:acceptor blends.

Electron affinity. To study the effect of the electron affinity on the CTC formation, we investigated the $\alpha(C_a)$ dependencies for MEH-PPV blends with different fluorene acceptors: **TNFon**, **HeptCTNFon**, **BuCDDNF**, **DTNF**, **AmCDTNF**. All these blends demonstrated pronounced threshold-like complexation behavior (Fig. 4a–e). The threshold acceptor concentration, C_a^t , that corresponds to the beginning of intensive polymer complexation varies by about 50 times for different acceptors (10 mM for **BuCDDNF** and 0.2 mM for **AmCDTNF**, see Fig. 4). It is logical to suppose that C_a^t is determined by the CTC binding energy, and the lower this concentration, the larger the CTC binding energy. Indeed, the lowest C_a^t is observed in the blends with strong acceptors **AmCDTNF** and **DTNF** (cf. LUMO energies of acceptors, Table 1 and Figs. S1 and S2 in SI). This is in line with the Mulliken model, where the CTC binding energy depends on the inverse effective HOMO–LUMO gap between the interacting donor and acceptor, and it increases with increasing the acceptor electron affinity.²²

Donor and Acceptor Molecular Skeletons. To address the effect of the acceptor and donor molecular skeletons, we also studied MEH-PPV blends with non-fluorene acceptors (**DNAQ** and **TCNQ**), and P3HT:**TNFon** blend. Compared to the MEH-PPV:fluorene acceptor blends, the MEH-PPV:**DNAQ** blend showed weaker inflexion in the $\alpha(C_a)$ dependence (Fig. 4f). Moreover, the MEH-PPV:**TCNQ** and P3HT:**TNFon** blends demonstrated virtually linear $\alpha(C_a)$ dependencies (Fig. 4g,h). These linear dependencies can be attributed to either the absence of a threshold or to a high threshold acceptor concentration that cannot be reached experimentally because of the limited acceptor solubility. Note that the CTC concentration, C_{CTC} , increases linearly below the threshold in

the MEH-PPV:**TNFon** blend.²⁰ Unfortunately, we cannot certainly distinguish between these two cases as the CTC molar absorption coefficients and hence the absolute CTC concentration are unknown (see below). However, in P3HT:**TNFon** blend, the used acceptor concentration was very high (up to 100 mM) to observe the charge-transfer band, so we assume that the threshold should be observed if it would exist. As it was not observed at all (α depends on C_a near linearly or even with slightly negative curvature as seen in Fig. 4h and Fig. S5h in SI), we conclude that complexation in P3HT:**TNFon** blends does not follow the threshold-like behavior. So, the threshold-like complexation apparently is not an universal property for all CP:acceptor blends but occurs only in some of them, and the molecular skeletons (geometry, rigidity, electronic structure etc.) of both polymer donor and low-molecular acceptor are important.

Solvent Effect. To investigate the effect of solvent on the CTC formation, we studied the $\alpha(C_a)$ dependencies for MEH-PPV:**TNFon** blends in aromatic and non-aromatic solvents of different polarity and polarizability (Table S1 in SI): chlorobenzene (CB), o-dichlorobenzene (DCB), toluene (Tol), chloroform (CF), and tetrahydrofuran (THF). Fig. 5 shows blend absorption *versus* the acceptor concentration in these solvents. In all the solvents, the S-shape dependencies were observed in accordance to NE model, and the threshold acceptor concentrations (C_a^t) were significantly different (Fig. 5a). While the solvent effect on C_a^t is apparent, it is not well described quantitatively by any solvent parameter (dipole moments, Kirkwood and Lippert-Mataga parameters, etc., see Table S1 in SI). It seems understandable, because in contrast to small molecular systems, polymer-acceptor-solvent interaction is a more complex process. Even for pristine MEH-PPV in solution, its conformation can change from a collapsed state to extended and planar coils depending on the solvent properties.⁴³⁻⁴⁷ To visualize the solvent effect on the slopes of $\alpha(C_a)$ dependencies, we have normalized the $\alpha(C_a)$ graphs in Fig. 5a to both C_a^t and absorbance maxima (Fig. 5b; see also the data normalized to C_a^t only in Fig. S8, SI). While the full plateaus have not been achieved in all the solvents due to limited solubility, differences in the slopes are clearly seen for different solvents (Fig. 5b and Fig. S8). However, the slopes do not correlate with any solvent parameter as well (e.g. see Table S1 in SI), and this could also be attributed to the complexity of the polymer-acceptor-solvent system.

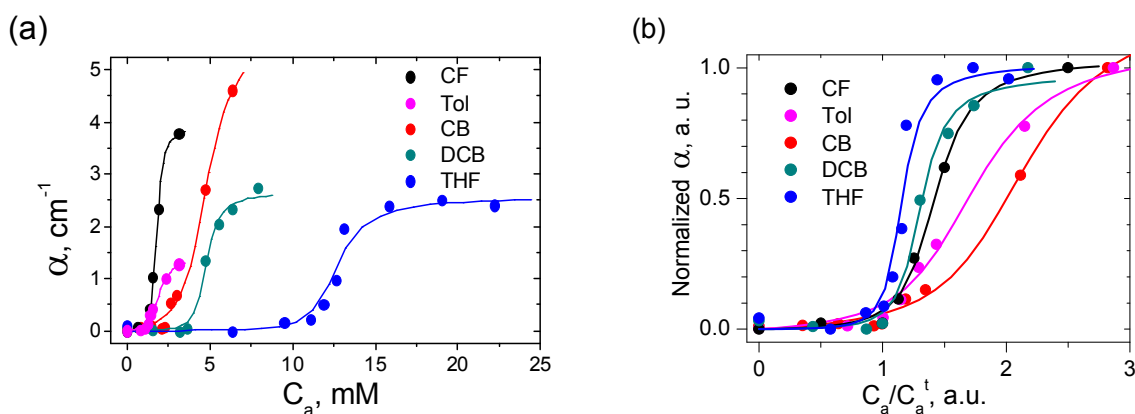


Fig. 5. CTC absorption at 630 nm *versus* the acceptor concentration for MEH-PPV:**TNFon** blends in different solvents. The MEH-PPV concentration was 7.68 mM for CB and 1.91 mM for other solvents (the absorption coefficients for CB were divided by four). The cuvette path length was 0.10 mm. Points are the experimental data, lines are the NE model fits. (a) As-measured data; (b) the data normalized to C_a^t (x axis) and absorption coefficient at maximum acceptor concentration (y axis).

Polymer Concentration. To address the impact of the polymer concentration and correspondingly the inter-macromolecular interactions (i.e. that between the segments of different polymer chains) on the CTC formation, we compared the $\alpha(C_a)$ dependencies for two series of MEH-PPV:TNFon blends with significantly different polymer concentrations: 2 g/L (7.68 mM) and 0.01 g/L (38.4 μ M). The lower concentration corresponds to the diluted regime, where the polymer coils are far from each other. The higher concentration corresponds to the semi-diluted regime, where the polymer coils overlap.²⁴ The 200-fold polymer concentration increase from the diluted to semi-diluted regime significantly increases the number of inter-macromolecular contacts and hence should considerably enhance the inter-macromolecular interactions. Therefore, if the inter-macromolecular interactions are of key importance for the CTC formation and/or are responsible for threshold-like CP complexation, the corresponding effects would be more pronounced for more concentrated solution and weaker for the dilute one. Fig. 6 represents the obtained average number of CTC at the polymer chain, $\langle N \rangle$, as a function of the acceptor concentration for the two blends. $\langle N \rangle$ was calculated from the absorption data:

$$\langle N \rangle = \frac{C_{CTC}}{C_d / N_d} = \frac{\alpha \cdot N_d}{\varepsilon \cdot C_d} = \frac{\alpha}{\varepsilon \cdot C_d} \cdot \frac{M_n}{\mu}, \quad (1)$$

where C_d is a concentration of the monomer repeat units, N_d is a number of repeat units per polymer chain, and hence C_d/N_d is a concentration of polymer chains, M_n is a number average molar mass of the polymer, μ is a molar mass of the repeat unit, and ε is a CTC molar extinction. For ε , an estimation for MEH-PPV:TNFon CTC in CB from Ref. 6 ($\varepsilon = 12700 \text{ M}^{-1}\text{cm}^{-1}$) was used. Fig. 6 demonstrates that the $\langle N \rangle(C_a)$ dependencies for the blends for the substantially different donor concentrations are very close indicating that CTC formation is not sensitive to the polymer concentration and hence to the inter-macromolecular interactions. Therefore, we conclude that the latter do not play a major role in the CP complexation and the polymer complexation operates mainly at individual polymer chains. Notably, the curve shape and the threshold acceptor concentration (Fig. 6) are not affected by the polymer concentration indicating that the threshold-like behavior does not originate from the inter-macromolecular interactions.

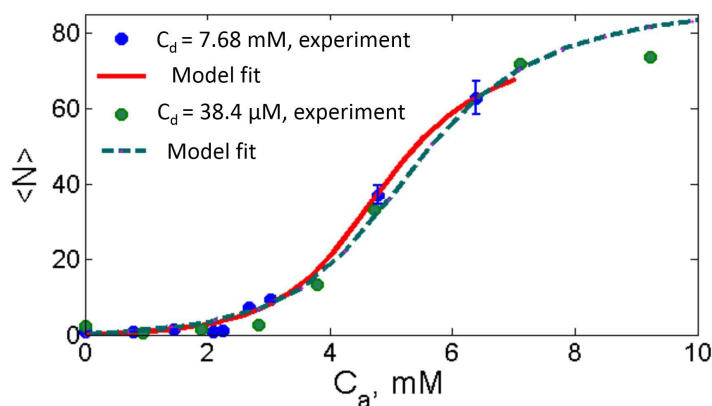


Fig. 6. Average CTC numbers per polymer chain *versus* acceptor concentration for MEH-PPV:TNFon blends in chlorobenzene at polymer concentration of 2 g/L (7.68 mM) and 0.01 g/L (38.4 μ M). Dots are the experimental data, lines are the fits by the NE model.

Discussion

Neighbor Effect Model. To fit the experimental data, we will use the NE model¹⁹ briefly outlined in this section. According to the model, the CTC is formed between an acceptor molecule and the polymer segment expanded to several repeat units of the polymer main chain. The binding energy of a CTC is E_0 if both the neighboring polymer segments are uncomplexed (Fig. 1). It increases by E_{NE} if one of them is complexed and by $2E_{NE}$ if both are complexed. Note that, according to the binding energy definition, E_{NE} corresponds to the total energy gain for the whole system if two CTCs are formed nearby, i.e., the effective binding energy per acceptor molecule increases by $E_{NE}/2$. The model gives a dependence of the average CTC number per chain, $\langle N \rangle$, on the acceptor concentration, C_a . Since $\langle N \rangle$ is proportional to the CTC concentration in solution (Eq. (1)), the model gives the $C_{CTC}(C_a)$ dependence as well. The NE model is analytical and has three fitting parameters: (i) the number of sites available for the CTC formation per chain (L), (ii) the binding energy of an isolated CTC (E_0), and (iii) the energy gain for two CTCs formed nearby (E_{NE}). In this work, we use a simplified version of the NE model that neglects a decrease in the acceptor concentration due to its involvement in CTCs; the reasons for this will be specified below. Interestingly, in this simplified NE model, the $\langle N \rangle(C_a)$ curve shape is determined solely by E_{NE} and does not depend on L or E_0 within a reasonable range of the parameters.¹⁹ The two remaining model parameters are responsible only for axes scaling: L determines the “y” scale, and the “x” scale is determined by $E_0 + E_{NE}$. Fig. 7 shows the average share of complexed segments $\langle N \rangle/L$ as a function of the acceptor concentration for different E_{NE} values (the “x” scale is normalized to the acceptor concentration at which the half of the chain is complexed). One can see that the larger the E_{NE} , the steeper the threshold. Therefore, since $\langle N \rangle$ is proportional to C_{CTC} , the shape of the experimental $C_{CTC}(C_a)$ curve provides information about the NE strength in terms of E_{NE} . The threshold acceptor concentration C_a^t that corresponds to the beginning of the CTC concentration increase is defined in the modeling as shown in Fig. 7. Since the scale of “x” axis (acceptor concentration) is determined by $E_0 + E_{NE}$, the threshold acceptor concentration is also associated with $E_0 + E_{NE}$. As a result, analysis of the $C_{CTC}(C_a)$ data with the NE model allows us to estimate the isolated CTC binding energy and NE in various CP:acceptor blends.

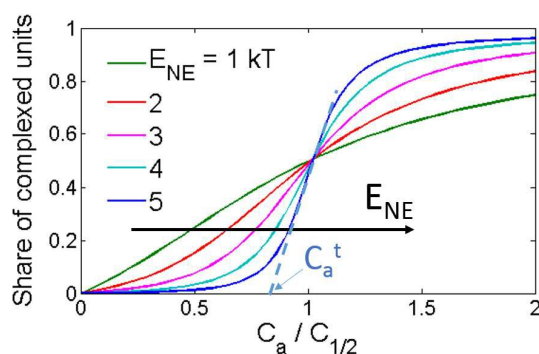


Fig. 7. Share of the complexed repeat units *versus* acceptor concentration for various E_{NE} according to the NE model. The acceptor concentration is normalized to the concentration at which half of the monomer units of the polymer is complexed, $C_{1/2}$. C_a^t shows the threshold acceptor concentration for the curve with $E_{NE}=5$ kT.

Fitting the Data with NE Model. The model described above deals with the CTC concentration as a function of the acceptor concentration, $C_{CTC}(C_a)$. We can obtain $C_{CTC}(C_a)$ from the measured CTC absorption, α : $C_{CTC}(C_a) = \alpha(C_a)/\epsilon$, where ϵ is the CTC molar extinction. However, the CTC molar extinction was determined only for one blend, i.e. MEH-PPV:**TNFon**,⁶ and it is unknown for the blends of other acceptors studied in this work. Unfortunately, Benesi-Hildebrand and related linear

regression methods,^{41, 48,49} used for determination of CTC molar extinction coefficients that usually applied for small-molecule CTC and the method proposed in Ref. 6 for threshold-like $\alpha(C_a)$ are inapplicable. The first methods are inappropriate since $\alpha(C_a)$ in our polymer donor–molecular acceptor mixtures is not described by the standard association constant model.⁶ The latter method assumes that the ratios between the CTC molar extinction at some wavelength and the polymer molar extinction are the same in solution and in film. However, this assumption is not solid for most of our blends as we found that the film spectra differ significantly from that in solutions (see Fig. S6b in SI). Accordingly, for using the assumption of constant ratio between the CTC and polymer molar extinctions, one needs to separate the polymer and the CTC absorptions in the solution and film spectra. Because this is not an easy task (if possible at all), we considered the CTC extinction as an additional fitting parameter and applied the model directly to the experimental $\alpha(C_a)$ dependencies. Doing that, we supposed that the CTC extinction for a pair MEH-PPV:acceptor is weakly affected by the number of acceptor molecules (one, two or more) involved in CTC on the same polymer chain. This seems reasonable taking into account that variations in extinctions of low-molecular π – π CTC are not high. Thus, for CTC of various fluorene acceptors with low-molecular donors, e.g. anthracene,⁴¹ N-ethylcarbazole,⁴⁹ or N-propylcarbazole,³⁹ in different solvents (1,2-dichloroethane, dioxane), the molar extinction coefficients vary from 800 to 1900 M⁻¹ cm⁻¹ only, whereas variations in the equilibrium constants is much higher, from 1.07 to 12.1 M⁻¹. In line with this, in MEH-PPV:**TNFO**n blends, the CTC molar extinctions are independent of the CTC concentration.⁶ For non-complicated molecular complexation of D + A = CTC, an isobestic point is generally observed evidencing only two spectroscopically resolvable species. Isobestic points are also observed for some our complexes (see Fig. 3a,b and Fig. S4a,b in SI), although not always for full concentration regions. This could be understood taking into account flexibility of MEH-PPV backbone and its conformation changes induced by an environment (e.g. by solvent as discussed above, but the same effect is one would expect on CTC formation). This would result in an intensity change of the main MEH-PPV absorption band and its bathochromic shift due to planarization of the polymer backbone^{43-47,50} so that the isobestic point can be smeared. Yet, an observation of relatively clear isobestic points in many cases of MEH-PPV:acceptor complexations allow us to consider that just one type of CTC would be sufficient to analyze the data in the NE model.

We use here the simplified NE model (described above) as it is much less sensitive to the CTC concentration and unknown CTC molar extinction. As mentioned above, the shape of the $C_{CTC}(C_a)$ curve and its x-axis scaling are practically independent of the chain length L in the simplified NE model. In other words, the average share of the complexed segments, $\eta(C_a) = \langle N \rangle / L$, is independent of L for its reasonable values and has only two parameters, E_0 and E_{NE} . As a result, the CTC concentration and therefore the CTC absorption are nearly linearly proportional to L (eq. 2):

$$\alpha = \varepsilon \cdot C_{CTC} = \varepsilon \cdot \frac{C_d}{N_d} \cdot \langle N \rangle \approx \varepsilon \cdot L \cdot \frac{C_d}{N_d} \cdot \eta(E_0, E_{NE}; C_a) \quad (2)$$

Therefore, in the simplified NE model, the $\alpha(C_a)$ dependence has three fitting parameters: E_0 , E_{NE} , and $(\varepsilon \cdot L)$, and the model remains three-parametrical. We will focus here on evaluation of the CTC energetics, E_0 and E_{NE} , without discussing the third fitting parameter, $\varepsilon \cdot L$. Note that a possible inaccuracy in $\varepsilon \cdot L$ does not influence the E_0 and E_{NE} best-fit estimates in the simplified NE model.¹⁹

The results of the data fitting are presented in Fig. 4 by solid lines. The model fits well the experimental data providing reasonable estimates for E_0 in the range of 2–6 kT and for E_{NE} in the range of 0–4 kT. The E_0 values are of the same order as the CTC binding energy in small-molecule donor:acceptor blends.⁵¹ Remarkably, in some of the blends, E_{NE} can be close to or even larger than E_0 , as we have earlier found for MEH-PPV:**TNFO**n blends.¹⁹ Therefore, the NE can play a key role

in the CTC formation in CP:acceptor blends. On the other hand, for MEH-PPV:TCNQ and P3HT:TNFon blends, the experimental data do not show any S-type features and can be fitted well by a straight line (Fig. 4g,h). Moreover, some degree of negative curvature is observed for P3HT:TNFon blends at high acceptor concentrations (Fig. S5g,h) so that the data can be fitted by common models for low-molecular CTCs. Therefore, assuming that a significant part of the CP chains is complexed (see above), we suggest a negligibly small NE (see Fig. 7) in MEH-PPV:TCNQ and P3HT:TNFon blends.

In the next section, using the results of $\alpha(C_a)$ fitting, we summarize how the isolated-CTC binding energy (E_0) and NE extra binding energy (E_{NE}) depend on the acceptor electron affinity, solvent, polymer concentration, and molecular skeletons of the donor and acceptor.

Factors Governing the CTC Binding Energy and the NE. The effect of the acceptor electron affinity on E_0 and E_{NE} was investigated in MEH-PPV blends with various fluorene acceptors. Fig. 8a demonstrates the best-fit values for E_0 versus the acceptor electron affinity, E_A . Note that the E_0 value (in combination with E_{NE}) relates to the threshold acceptor concentration C_a^t .¹⁹ The higher E_0 , the lower the C_a^t . The observed up to 50-fold difference in C_a^t corresponds to the variation of E_0 by several times (cf. BuCDDNF and AmCDTNF in Fig. 4 and Fig. 8a). One can see that E_0 has a trend of growth with increasing the acceptor electron affinity (at least for structurally related fluorene acceptors), in accordance with the Mulliken model.²²

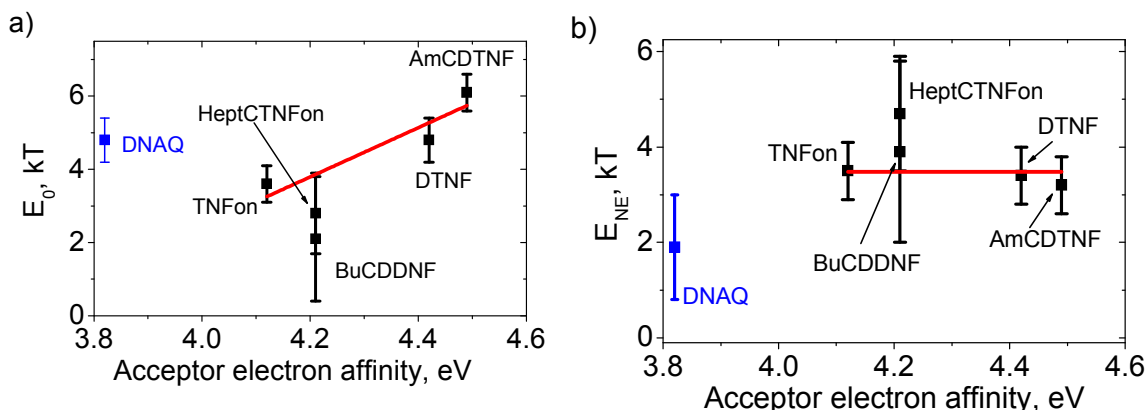


Fig. 8. Binding energy of an isolated CTC (a) and NE energy gain (b) obtained from the fits of the absorption data (Fig. 4). The lines are a guide to the eye for the fluorene acceptors.

Fig. 8b presents the calculated E_{NE} values versus the electron affinity of the studied acceptors. Unexpectedly, all MEH-PPV:fluorene acceptor blends show near the same E_{NE} values within the fitting error. This corresponds to a similar $\alpha(C_a)$ shape for these blends (Fig. 4a-e) as the latter is determined solely by E_{NE} in the NE model (see above). Therefore, one can conclude that the NE does not depend on the acceptor electron affinity within the studied range.

As shown above, the $\alpha(C_a)$ dependencies for MEH-PPV:TNFon blends in various solvents differ by both curvature of the S-shaped profiles and threshold acceptor concentration, C_a^{thr} (Fig. 5). As a result, fitting the data with the NE model gives different E_0 and E_{NE} parameters. Fig. 9 shows the best-fit values of E_0 (panel a) and E_{NE} (panel b) in different solvents, obtained from the data in Fig. 5, versus the Lippert-Mataga index, Δf . While the regression is weak, the general trend is obvious: the larger the Δf , the lower the binding energy of an isolated CTC (E_0), and the higher the NE (E_{NE}).

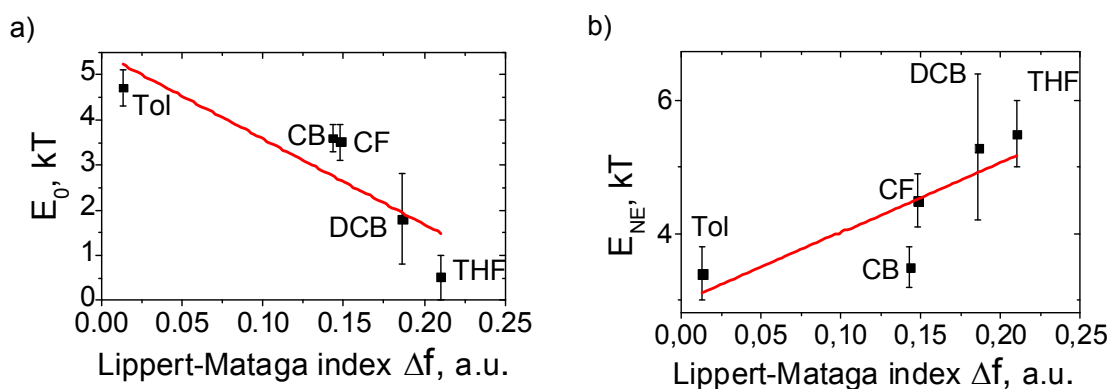


Fig. 9. E_0 and E_{NE} values in different solvents obtained from the data in Fig.5 versus Lippert-Mataga index $\Delta f = (\epsilon - 1)/(2\epsilon + 1) - (n^2 - 1)/(2n^2 + 1)$, where ϵ is the dielectric permittivity and n is the refraction index. The error bars are given at a confidence level of 80%. The lines are a guide to the eye.

The experimental data in Fig. 6 show that the $\alpha(C_d)$ curves practically coincide for MEH-PPV:**TNFon** blends with the donor concentration differing by 200 times. As a result, the best-fit parameter values for the blend with the low C_d ($E_0 = 3.6 \pm 0.8$ kT, $E_{NE} = 3.3 \pm 0.8$ kT) coincide within the experimental and fitting errors with those for the blend with the high C_d ($E_0 = 3.6 \pm 0.5$ kT, $E_{NE} = 3.5 \pm 0.6$ kT). This indicates that the CTC binding energy and the NE do not depend on the polymer concentration and hence on the inter-macromolecular interactions.

For MEH-PPV blends with **DNAQ** and **TCNQ**, whose conjugated molecular skeletons considerably differ from the fluorene ones, and P3HT:**TNFon** blends, in which the donor has a conjugated skeleton different from that of MEH-PPV, the $\alpha(C_d)$ shapes are significantly different from those of the MEH-PPV blends with fluorene acceptors (see Fig. 4). The NE model attributes this to much weaker NE in the former blends (see Fig. 7). For **DNAQ**, $E_{NE} = 1.9 \pm 1.1$ kT, which is about 1.5 times lower than the average E_{NE} for the MEH-PPV blends with fluorene acceptors. This indicates that the NE is sensitive to the size and the shape of conjugated molecular core of an acceptor. For MEH-PPV:**TCNQ** and P3HT:**TNFon** blends, the linear $\alpha(C_d)$ dependencies can be treated as evidence for weak or absent NE in these blends assuming that the majority of polymer conjugated segments is complexed at the maximum acceptor concentration (see above). Therefore, we conclude that the NE is very sensitive to the shape and size of conjugated molecular cores of both the donor and acceptor. The isolated CTC binding energy for MEH-PPV:**DNAQ** blend is $E_0 = 5.0 \pm 0.7$ kT, which is relatively large as compared to the values for the MEH-PPV:fluorene acceptors, taking into account the lower electron affinity of **DNAQ**. This underlines that E_0 is also sensitive to the structures of the donor and acceptor conjugated molecular cores.

Table 2 summarizes our findings on the CTC energetics in the studied polymer:acceptor blends. Both energies E_0 and E_{NE} are insensitive to the polymer concentration indicating that the complexation can operate at a single chain and the NE is an intra-macromolecular effect. While the isolated CTC binding energy E_0 depends on the solvent quality, acceptor electron affinity, and molecular skeletons of the donor-acceptor pair, only two of the investigated factors, namely the type of donor and acceptor conjugated molecular skeletons and the solvent, determine the NE energy gain in CP complexation, E_{NE} . In the next section, we rationalize and discuss possible microscopic mechanism that could be responsible for the NE.

Table 2. Factors affecting E_0 and E_{NE} in CP complexation with small molecule acceptors according to the NE model.

Factor	Sensitivity of the NE model parameters	
	E_0	E_{NE}
Polymer concentration	No	No
Solvent	Yes	Yes
Electron affinity	Yes	No
Conjugated molecular skeleton	Yes	Yes

Microscopic Picture of the Neighbor Effect

The current NE model is one-dimensional (1D) and does not take into account directly any 3D effects as, e.g., changes in the conformational state of the polymer and its aggregation, which can affect the CTC formation and seem to be essential in CP CTCs. Nevertheless, the NE model can be a starting point to analyze the microscopic mechanism underlying the CTC formation in CPs. We suggest that the NE stems from chain planarization upon CTC formation. While only a few repeat units of the polymer can directly contact a short acceptor molecule in a CTC (via overlapping of the polymer and acceptor orbitals), the electronic and conformational state of the rest of the conjugated segment (ca. 5–7 units in MEH-PPV^{7, 52, 53}) is also affected by complexation.^{16,54} Specifically, using Raman spectroscopy, the CTC formation was shown to increase the chain planarity and torsional rigidity^{16,55} that can be explained by more quinoid character of the complexed polymer chain, which is stiffer and more planar. The NE can then be attributed to the two mechanisms: steric and/or electronic. First, the complexed polymer segment is more planar and therefore sterically more favorable for the subsequent CTC formation with a planar acceptor as illustrated in Fig. 9a (steric mechanism). Indeed, the planarized chain can provide larger orbital overlapping with the π -electron conjugated orbitals of planar acceptor increasing the electronic coupling between the donor HOMO and acceptor LUMO, H_{DA} , and hence the CTC binding energy. In addition, because of more quinoid character, the effective conjugation length of the complexed chain is less limited by the thermally activated torsional deformations,⁵⁵ and hence it is longer than that of the free (uncomplexed) chain. Fig. 9b (left) illustrates an increase of the effective conjugation length of a CP after CTC formation with an acceptor molecule. The increased conjugation length results in a higher/lower local HOMO/LUMO level as illustrated in Fig. 9b (right). Note that the increase in the HOMO energy due to planarization can overwhelm its decrease due to CTC formation. According to the Mulliken model, the lower the energy difference between the donor HOMO and the acceptor LUMO, the larger the CTC binding energy (electronic mechanism).²² Therefore, the latter can be higher for CTCs formed at a conjugated segment that already has a CTC, i.e., at the repeat units adjacent to the complexed ones.

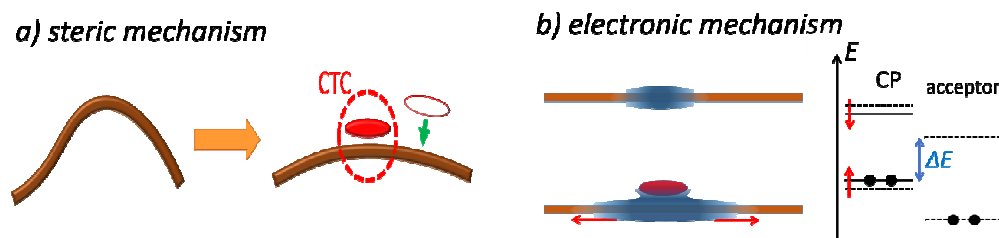


Fig. 9. Illustration of the two mechanisms of the NE according to the chain planarization hypothesis. a) Steric mechanism. After the first CTC formation, the chain becomes more planar making formation of successive CTCs more favorable since the planarized chain can provide larger overlapping with the acceptor π -electron conjugated system. b) Electronic mechanism. (left) Increasing the effective conjugation length of a CP upon CTC formation. Top: a free CP chain (brown) with the effective conjugation length depicted by a blue region. Bottom: a CP chain complexed with an acceptor molecule (red oval) has an extended effective conjugation length

(indicated by red arrows). (right) Frontier molecular orbital energies of a pair of non-interacting polymer conjugated segment and acceptor molecule (dashed), and that of a complexed polymer conjugated segment (solid). ΔE stands for effective HOMO(D) – LUMO(A) gap. Red arrows illustrate shifts of the frontier orbital energies of the polymer conjugated segment as a result of CTC formation.

We suggest that the chain planarization upon CTC formation originates from the more quinoid character of the complexed chain, and hence is determined by the degree of charge transfer in a CTC. In the Mulliken model, the degree of charge transfer is determined by two factors: donor-acceptor electronic coupling, H_{DA} , and the effective HOMO(D)-LUMO(A) gap. The observed strong dependence of the NE on the donor and acceptor molecular structure is in complete accordance with the chain planarization hypothesis since the molecular structures determine H_{DA} . However, we observed that the EA does not affect the NE in CP complexation that seems at some contradiction with the hypothesis. In order to reconcile the data with the hypothesis, we assume that with EA increase the charge density on acceptor shifts from the fluorene core to the substituents, and this shift in turn can decrease H_{DA} and limit the net charge transfer degree making the latter less sensible to EA . With this assumption, the chain planarization hypothesis appears to agree with the experimental data; however, the more detailed analysis of the suggested NE mechanism is beyond the scope of the current study. Consequently, we suggest that the chain planarization hypothesis describes a possible microscopic mechanism of the NE in the CP complexation. One can expect that the NE will be strong in blends where the CTC formation substantially increases the chain planarity. This would require a planar acceptor with an extended π -system (and reasonably low LUMO) for strong orbital overlap and pronounced π - π CT interaction with polymer donor moieties. In contrast, the NE is expected to be negligible in polymer:fullerene blends, e.g. P3HT:PCBM or MEH-PPV:C₆₀, as well as in blends with very small acceptors (e.g. tetracyanoethylene, TCNE, or 2,3-dichloro-5,6-dicyanobenzoquinone, DDQ). Indeed, as the fullerene molecules are not planar, and the size of the conjugated system of very small acceptors is much smaller than the polymer conjugated length, overlapping between the polymer HOMOs and the acceptor LUMOs should be very weak, and the chain planarity and electronic structure will be slightly affected by the CTC formation thus resulting in a weak NE. We also argue that the NE should operate stronger for CPs with semi-flexible chains where the complexation could noticeably increase the chain planarity.

Additional contribution to the NE can stem from cross-chain π - π aggregation that was initially suggested in Ref. 6. Following this suggestion, a CTC could be a link between two distant segments of a single polymer chain. The formation of the CTC-link and subsequent planarization of the two segments would result in approaching their monomer units adjacent to the link and an increased probability for successive CTC-links to be formed nearby the first one (Fig. 3 in Ref. 6). Note that the NE was found to be independent of the polymer concentration (the latter was changed by 200 times, Fig. 6), which determines the number of interchain contacts. Therefore, interaction of different macromolecules does not affect the NE, which operates within an individual macromolecule.

A strong NE should result in high inhomogeneity of the CTC distribution over the chains so that large CTC aggregates can be formed.¹⁹ These aggregates can be inherited in film, where microscopic ordered domains of the complexed polymer were observed.²¹ The morphology and photophysics is completely different in the domains of neat and complexed CP.² The NE strength can determine the size of the CTC domains and thus could be used to control the film morphology. Moreover, blend self-organization as a result of the NE can be a route to overcome the intrinsic disorder of CPs and result in improved performance of polymer electronic devices. Therefore, search for CP blends with strong NE deserves further studies.

Conclusions

We have found that in a number of MEH-PPV blends with various molecular π -electron acceptors, the charge-transfer complex (CTC) absorption shows threshold-like dependence on the acceptor concentration, while in the other MEH-PPV and P3HT-based blends this dependence resembles that typical for small molecule donor-acceptor blends. The optical absorption data were analyzed within a model that takes into account the neighbor effect (NE) in conjugated polymer (CP) complexation. We conclude that the NE can play a key role in this complexation. The binding energy of an isolated CTC was found to be sensitive to the acceptor electron affinity, the solvent, and the type of donor and acceptor conjugated molecular skeletons, while the NE depends only on the skeletons and the solvent. We suggest that the NE in CP complexation operates within an individual macromolecule and stems from the planarization of the conjugated chain upon CTC formation. We expect that the NE should be weak in CP blends with non-planar acceptors (e.g. fullerene derivatives), acceptors with relatively small π -delocalized orbitals, and blends with relatively rigid CPs, whereas it is expected to be more pronounced in blends of semi-flexible CPs with acceptors having an extended π -conjugated system suitable for efficient orbital overlap with polymer donor moieties.

Supporting Information

Synthesis and characterization of studied fluorene acceptors with respective ^1H and ^{13}C NMR spectra, mass-spectra, infrared spectra, melting points and elemental analyses. Additional optical absorption data for polymer:acceptor blends in solutions and in films. DFT calculations for fluorene acceptors and correlations between computational DFT data and CV experiments.

Acknowledgement

This work was supported by Russian Foundation of Basic Research (projects #13-02-01313 and 13-03-12472). This work was done using equipment purchased under the Lomonosov Moscow State University Program of Development. S.K. thanks to Bangor University for 125 Anniversary PhD Scholarship.

References

1. A. A. Bakulin, S. G. Elizarov, A. N. Khodarev, D. S. Martyanov, I. V. Golovnin, D. Y. Paraschuk, M. M. Triebel, I. V. Tolstov, E. L. Frankevich, S. A. Arnautov and E. M. Nechvolodova, *Synth. Met.*, 2004, **147**, 221-225.
2. A. Y. Sosorev and D. Y. Paraschuk, *Isr. J. Chem.*, 2014, **54**, 650-673.
3. D. Y. Paraschuk, S. G. Elizarov, A. N. Khodarev, A. N. Shchegolikhin, S. A. Arnautov and E. M. Nechvolodova, *JETP Letters*, 2005, **81**, 467-470.
4. A. A. Bakulin, M. S. Pshenichnikov, P. H. M. van Loosdrecht, I. V. Golovnin and D. Y. Paraschuk, in *Physics of nanostructured solar cells*, eds. V. Badescu and M. Paulescu, Nova Science Publishers, 2010, pp. 463-504.
5. A. A. Bakulin, A. N. Khodarev, D. S. Martyanov, S. G. Elizarov, I. V. Golovnin, D. Y. Paraschuk, S. A. Arnautov and E. M. Nechvolodova, *Dokl. Chem.*, 2004, **398**, 204-206.
6. O. D. Parashchuk, V. V. Bruevich and D. Y. Paraschuk, *Phys. Chem. Chem. Phys.*, 2010, **12**, 6021 - 6026.

7. A. A. Bakulin, D. S. Martyanov, D. Y. Paraschuk, M. S. Pshenichnikov and P. H. M. van Loosdrecht, *J. Phys. Chem. B.*, 2008, **112**, 13730-13737.
8. A. E. Ozimova, V. V. Bruevich, T. Dittrich and D. Y. Paraschuk, *Macromol. Symp.*, 2010, **296**, 138-143.
9. A. A. Bakulin, S. A. Zapunidy, M. S. Pshenichnikov, P. H. M. van Loosdrecht and D. Y. Paraschuk, *Phys. Chem. Chem. Phys.*, 2009, **11**, 7324-7330.
10. J. J. Benson-Smith, L. Goris, K. Vandewal, K. Haenen, J. V. Manca, D. Vanderzande, D. D. C. Bradley and J. Nelson, *Adv. Funct. Mater.*, 2007, **17**, 451-457.
11. T. Drori, C. X. Sheng, A. Ndobe, S. Singh, J. Holt and Z. V. Vardeny, *Phys. Rev. Lett.*, 2008, **101**, 037401.
12. T. Drori, J. Holt and Z. V. Vardeny, *Phys. Rev. B*, 2010, **82**, 075207.
13. C. Deibel, T. Strobel and V. Dyakonov, *Adv. Mater.*, 2010, **22**, 4097-4111.
14. T. M. Clarke and J. R. Durrant, *Chem. Rev.*, 2010, **110**, 6736-6767.
15. K. Vandewal, K. Tvingstedt and O. Inganäs, in *Semiconductors and Semimetals*, eds. M. T. Uli Würfel and R. W. Eicke, Elsevier, 2011, vol. Volume 85, pp. 261-295.
16. V. V. Bruevich, T. S. Makhmutov, S. G. Elizarov, E. M. Nechvolodova and D. Y. Paraschuk, *J. Chem. Phys.*, 2007, **127**, 104905.
17. S. G. Elizarov, A. E. Ozimova, D. Y. Paraschuk, S. A. Arnautov and E. M. Nechvolodova, *Proc. SPIE*, 2006, **6257**, 293-302.
18. I. V. Golovnin, A. A. Bakulin, S. A. Zapunidy, E. M. Nechvolodova and D. Y. Paraschuk, *Appl. Phys. Lett.*, 2008, **92**, 243311.
19. A. Sosorev and S. Zapunidi, *J. Phys. Chem. B*, 2013, **117**, 10913-10919.
20. A. Y. Sosorev, O. D. Parashchuk, S. A. Zapunidi, G. S. Kashtanov and D. Y. Paraschuk, *J. Phys. Chem. C*, 2013, **117**, 6972-6978.
21. O. D. Parashchuk, S. Grigorian, E. E. Levin, V. V. Bruevich, K. Bukunov, I. V. Golovnin, T. Dittrich, K. A. Dembo, V. V. Volkov and D. Y. Paraschuk, *J. Phys. Chem. Lett.*, 2013, **4**, 1298-1303.
22. R. S. Mulliken, *J. Am. Chem. Soc.*, 1952, **74**, 811-824.
23. D. Veldman, S. C. J. Meskers and R. A. J. Janssen, *Adv. Funct. Mater.*, 2009, **19**, 1939-1948.
24. O. D. Parashchuk, T. V. Laptinskaya and D. Y. Paraschuk, *Phys. Chem. Chem. Phys.*, 2011, **13**, 3775-3781.
25. V. G. Pavelyev, O. D. Parashchuk, M. Krompiec, T. V. Orekhova, I. F. Peregichka, P. H. M. van Loosdrecht, D. Y. Paraschuk and M. S. Pshenichnikov, *J. Phys. Chem. C*, 2014, **118**, 30291-30301.
26. Gaussian 09, Revision A02, 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, Inc., Wallingford CT, 2009.
27. A. D. Becke, *Phys. Rev. A*, 1988, **38**, 3098-3100.
28. A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648-5652.
29. C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B*, 1988, **37**, 785-789.
30. S. Miertus, E. Scrocco and J. Tomasi, *Chem. Phys.*, 1981, **55**, 117-129.

31. M. S. Newman, T. B. Patrick, R. S. Darlak and E. A. Zuech, *J. Org. Chem.*, 1969, **34**, 1904-1906.
32. E. H. Huntress, K. Pfister and K. H. T. Pfister, *J. Amer. Chem. Soc.*, 1942, **64**, 2845-2849.
33. G. I. Migachev, N. G. Grekhova and N. S. Dokunikhin, *Zh. Org. Khim. (Russ. J. Org. Chem.)* 1978, **14**, 2380—2387 (in Russian).
34. Hinderer, H. E.; Smith, R. W., US patent 3,467,623, 1969.
35. T. Sulzberg and R. J. Cotter, *J. Org. Chem.*, 1970, **35**, 2762-2769.
36. T. K. Mukherje and L. A. Levasseu, *J. Org. Chem.*, 1965, **30**, 644-646.
37. T. K. Mukherje, *J. Phys. Chem.*, 1966, **70**, 3848-3852.
38. I. F. Perepichka, A. F. Popov, T. V. Orekhova, M. R. Bryce, A. N. Vdovichenko, A. S. Batsanov, L. M. Goldenberg, J. A. K. Howard, N. I. Sokolov and J. L. Megson, *J. Chem. Soc., Perkin Trans. 2*, 1996, 2453-2469.
39. I. F. Perepichka, L. G. Kuz'mina, D. F. Perepichka, M. R. Bryce, L. M. Goldenberg, A. F. Popov and J. A. K. Howard, *J. Org. Chem.*, 1998, **63**, 6484-6493.
40. I. F. Perepichka, A. F. Popov, T. V. Orekhova, M. R. Bryce, A. M. Andrievskii, A. S. Batsanov, J. A. K. Howard and N. I. Sokolov, *J. Org. Chem.*, 2000, **65**, 3053-3063.
41. D. D. Mysyk, I. F. Perepichka and N. I. Sokolov, *J. Chem. Soc., Perkin Trans. 2*, 1997, 537-546.
42. P. Panda, D. Veldman, J. Sweelssen, J. J. A. M. Bastiaansen, B. M. W. Langeveld-Voss and S. C. J. Meskers, *J. Phys. Chem. B*, 2007, **111**, 5076-5081.
43. T.-Q. Nguyen, V. Doan and B. J. Schwartz, *J. Chem. Physics*, 1999, **110**, 4068-4078.
44. D. Wang, Y. Yuan, Y. Mardiyati, C. Bubeck and K. Koynov, *Macromolecules*, 2013, **46**, 6217-6224.
45. H. Yamagata, N. J. Hestand, F. C. Spano, A. Köhler, C. Scharsich, S. T. Hoffmann and H. Bässler, *J. Chem. Phys.*, 2013, **139**, 114903.
46. R. Traiphol, P. Sanguansat, T. Srihirin, T. Kerdcharoen and T. Osotchan, *Macromolecules*, 2006, **39**, 1165-1172.
47. D. Hu, J. Yu, K. Wong, B. Bagchi, P. J. Rossky and P. F. Barbara, *Nature*, 2000, **405**, 1030-1033.
48. H. A. Benesi and J. H. Hildebrand, *J. Am. Chem. Soc.*, 1949, **71**, 2703-2707.
49. S. A. Grebenyuk, I. F. Perepichka and A. F. Popov, *Spectrochim. Acta Part A: Mol. Spectrosc.*, 2002, **58**, 2913-2923.
50. Y. Ebihara and M. Vacha, *J. Phys. Chem., B*, 2008, **112**, 12575-12578.
51. R. S. Mulliken and W. B. Person, *J. Am. Chem. Soc.*, 1969, **91**, 3409-3413.
52. R. Chang, J. H. Hsu, W. S. Fann, K. K. Liang, C. H. Chang, M. Hayashi, J. Yu, S. H. Lin, E. C. Chang, K. R. Chuang and S. A. Chen, *Chem. Phys. Lett.*, 2000, **317**, 142-152.
53. T. E. Dykstra, E. Hennebicq, D. Beljonne, J. Gierschner, G. Claudio, E. R. Bittner, J. Knoester and G. D. Scholes, *J. Phys. Chem. B*, 2008, **113**, 656-667.
54. V. V. Bruevich, T. S. Makhmutov, S. G. Elizarov, E. M. Nechvolodova and D. Y. Paraschuk, *J. Exp. Theor. Phys.*, 2007, **105**, 469-478.
55. M. O. Osotov, V. V. Bruevich and D. Y. Paraschuk, *J. Chem. Phys.*, 2009, **131**, 094906.