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Synthesis of Polythiophene–Fullerene Hybrid Additives as Potential Compatibilizers of BHJ Active Layers

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Abstract: Perfluorophenyl functionalities have been introduced as side chain substituents onto regioregular poly(3-hexyl thiophene) (rr-P3HT), under various percentages. These functional groups were then converted to azides which were used to create polymeric hybrid materials with fullerene species, either C_{60} or C_{70}. The P3HT–fullerene hybrids thus formed were thereafter evaluated as potential compatibilizers of BHJ active layers comprising P3HT and fullerene based acceptors. Therefore, a systematic investigation of the optical and morphological properties of the purified polymer–fullerene hybrid materials was performed, via different complementary techniques. Additionally, P3HT:PC_{70}BM blends containing various percentages of the herein synthesized hybrid material comprising rr-P3HT and C_{70} were investigated via Transmission Electron Microscopy (TEM) in an effort to understand the effect of the hybrids as additives on the morphology and nanophase separation of this typically used active layer blend for OPVs.

Keywords: rr-polythiophenes; perfluorophenyl functionalization; fullerene-polymer hybrids; P3HT:PCBM blends; organic photovoltaics (OPVs)

1. Introduction

Bulk Heterojunction (BHJ) solar cell devices are widely investigated due to their potential to reach high efficiencies along with their potential of large area coverage and unique flexibility [1–3]. Current state of the art materials that form the bicontinuous interpenetrating network of the active layer are poly(3-hexylthiophene) (P3HT) and [6,6]-phenyl-C_{61}-butyric acid methyl ester (PCBM), providing power conversion efficiencies (PCE) of above 6%, because of their thermal stability [4], high electron mobility [5,6] and especially because it is the current system of choice for printable over large areas organic photovoltaics (OPVs) [7–10]. Among others, decisive parameters that affect the device performance are the materials comprising the active layer, the morphology of the blend as well as the device architecture. In the direction of materials development, new polymeric electron donating materials with complex architectures and lower energy levels [11–15] compared to P3HT, and new electron acceptors based on fullerene [16–18] or even fully organic and polymeric ones [19–21], have been reported. On the other hand, efforts to control the morphology [22,23] of the active blend have been made using thermal annealing or solvent treatment of the blend resulting in significantly higher PCEs [1,4,24–26]. Besides the above-mentioned manipulation of the active layer, another approach that has been followed is the modification of the semiconducting polymer with functional groups that may lead to more complex architectures either co-polymeric or hybrid ones. More specifically,
co-polymeric materials based on polythiophenes or other conjugated polymers that can interact with or bear fullerene species can provide better miscibility and tune the nanophase separation of the two plain blend components [27–33]. In some cases, the introduction of functional groups like hydroxyl or azide units, as side chain substituents or as end chain groups, not only controls but also stabilizes the morphology of the blended materials [34–38]. Another efficient way to control the morphology is the introduction into the active layer of small quantities of hybrid materials comprising the semiconducting polymer and a carbon nanostructure [39–48]. However, the synthesis of most semiconducting polymeric or copolymeric hybrids involves complicated and multistep synthetic procedures that affect the yield and the cost of the final materials. Moreover, in most such cases, a non-conjugated linkage between the semiconducting polymer and the carbon nanostructure has been employed preventing electronic interactions among the polymeric and fullerene parts of the hybrid materials. To overcome the above bottlenecks, new synthetic strategies involving the “direct” attachment of the two components, organic and fullerene ones, comprising the hybrid materials have been developed [49–54]. Working in this direction, we have recently reported a straightforward methodology for the development of hybrid semiconducting polymeric materials using the electron deficient perfluorophenyl functionality inserted either onto copolymeric materials or as end polymeric chain group [49,54]. Benefits of this methodology are the direct electronic interactions between the two components [50], the absence of extra aliphatic or other non-conjugated parts used as connection bridges which increase the volume of insulating material within the active layer as well as the use of the intermediate perfluorophenyl aromatic azides that are stable over time and ambient storage conditions.

In this present work, we focused on the introduction of perfluorophenyl functionalities as side chain substituents of regioregular-poly(3-hexyl thiophene) (rr-P3HT). Several ratios were attempted in order to investigate the influence of the side functionalization onto the properties of rr-P3HT. Thereafter, using our previously reported methodology [49,54], the perfluorophenyl functionalities were transformed into azides that performed a [3+2] cycloaddition reaction with C<sub>60</sub> or C<sub>70</sub> fullerene species to provide the desired hybrid materials. Extensive purification has been performed in all cases and the purified materials were characterized in respect to their optical and morphological properties. These particular hybrid materials are ideal candidates for the compatibilization and stabilization of P3HT:fullerene active layers used in BHJ polymer photovoltaics, since the introduction of the fullerene species directly along the polythiophene’s backbone ensures the hybrids’ incorporation in the nanophase created by the neat blend components potentially holding them in close proximity over time and temperature variations during device operation. In order to support the hybrids potentiality as compatibilizers, mixtures of typical P3HT:PC<sub>70</sub>BM active BHJ blends with various percentages of the C<sub>70</sub> based hybrid material were investigated in respect to the effect of the additive on the morphology of the blend.

2. Materials and Methods

2.1. Materials

rr-P3HT was synthesized according to literature procedure [55]. Fullerene C<sub>60</sub> (99.5%) was purchased from SES research. Carbon C<sub>70</sub> (99.5%) was kindly provided by Prof. Kyriakos Porfyrakis of the Oxford Carbon Nanomaterials Group, Department of Materials, University of Oxford, UK. Tetrahydrofuran was purchased from Aldrich (Sigma-Aldrich Chemie GmbH, Taufkirchen, Germany) and was distilled with benzophenone and metallic sodium (THF<sub>dry</sub>). All the other solvents and reagents were purchased from Aldrich and used without further purification unless otherwise stated.

2.2. Instrumentation

<sup>1</sup>H, <sup>19</sup>F, <sup>15</sup>N Nuclear Magnetic Resonance (NMR) spectra were recorded on a Bruker Advance (Bruker BioSpin GmbH, Magnet Division, Karlsruhe, Germany) DPX 400.13, 376.5 and 40.55 MHz spectrometer, respectively, in CDCl<sub>3</sub> containing TMS as internal standard.
Gel permeation chromatography (GPC) measurements were carried out using a Polymer Lab chromatographer (Agilent Technologies, Santa Clara, CA, USA) equipped with two PLgel 5 µm mixed columns and a UV detector, using CHCl₃ as eluent with a flow rate of 1 mL/min at 25 °C, calibrated versus polystyrene standards.

Attenuated Total Reflectance (ATR) spectra were recorded on a “Bruker Optics’ Alpha-P Diamond ATR Spectrometer of Bruker Optics GmbH” (Ettlingen, Germany).

Thermogravimetric analysis (TGA) was carried out on ~8 mg samples contained in alumina crucibles in a Labsys TM TG apparatus of Setaram (Caluire, France) under nitrogen and at a heating rate of 10 °C/min.

UV-Vis spectra were recorded using a Hitachi U-1800 spectrophotometer (Hitachi High-Technologies Europe GmbH, Mannheim, Germany). Continuous wave photoluminescence was measured on a Perkin Elmer LS50B spectrofluorometer (Waltham, MA, USA). All UV-Vis and PL measurements were performed in air using quartz cuvettes and flat substrates for the examination of solutions and films, respectively.

The Blend Preparation involved the mixing of stock solutions of P3HT (10 mg·mL⁻¹) and PC₇₀BM (10 mg·mL⁻¹) to prepare the P3HT:PC₇₀BM 1:1 w/w. Afterwards, 0.125 and 0.625 mL of the [P3HT₀.₆-(P3HT-5F)₀.₃-(P3HT-5F-N-C₇₀)₀.₁] (8 mg·mL⁻¹) were added to prepare the 20% and the 50% blends with the hybrid polymer, respectively.

Transmission electron microscopy (TEM) measurements were performed on a JEOL JEM2100 (Peabody, MA, USA) operating at 200 kV. Sample preparation for TEM examination of the [P3HT₀.₆-(P3HT-5F)₀.₄], [P3HT₀.₆-(P3HT-5F)₀.₂-(P3HT-5F-N-C₆₀)₀.₂], [P3HT₀.₆-(P3HT-5F)₀.₃-(P3HT-5F-N-C₇₀)₀.₁] involved the preparation of dilute solutions of the samples in o-DCB and filtration through a 0.45 µm filter. For the hybrids a drop of the solution was placed on 3 mm carbon coated copper grids (Electron Microscopy Sciences) and the samples were dried in air for 2 days. For the blends examination via TEM, a drop of the blends solution was placed on 3 mm carbon coated copper grids (Electron Microscopy Sciences) and after 30 s the solvent was removed with filter paper.

2.3. Synthetic Procedures

2.3.1. Side [P3HT₀.₆-(P3HT-Br)₀.₄] [56]

A 50 mL round bottom flask, equipped with a reflux condenser and a magnetic stirrer, was flamed under vacuum degassed and filled with argon. P3HT (500 mg, 3 mmol) was added and dissolved in 30 mL CHCl₃ and the system was degassed and flushed with argon again. The flask was covered with aluminum foil and NBS (214 mg, 1.2 mmol) was added in three portions. The mixture was stirred for 22 h at room temperature. Then, ~15 mL Na₂SO₃ 5% aqueous solution were added to the system and the mixture was extracted with CHCl₃ and deionized water. The organic phase was dried with anhydrous MgSO₄, filtrated and rotary evaporated. The obtained solid was dried under vacuum at 40 °C. Bromination ratio calculated from ¹H NMR: 40%, yield: 92% ¹H NMR (δH; CDCl₃; Me₄Si): 7.00(broad, 1H), 2.80(broad, 2H), 2.64(broad, 2H–Br substituted thiophene), 1.65(broad, 4H), 1.53–1.22(broad, 12H), 0.88(broad, 6H).

The same procedure was used for all other ratios.

2.3.2. Side [P3HT₀.₆-(P3HT-5F)₀.₄]

A 100 mL round bottom flask, equipped with a reflux condenser and a magnetic stirrer, was degassed (flamed under vacuum) and filled with argon. Side [P3HT₀.₆-(P3HT-Br)₀.₄] (400 mg, 0.81 mmol), pentafluorophenyl boronic acid (205 mg, 1 mmol) and Pd(PPh₃)₄ (30 mg, 0.02 mmol) were dissolved in 20 mL toluene and then Na₂CO₃ (1.20 mL, 2.42 mmol) in 2 M aqueous were added. The mixture was degassed and flushed with argon again and refluxed for 72 h. Then, the mixture was filtered in order to remove the catalyst and the solvent was rotary evaporated. The obtained solid was washed several times with MeOH and deionized water. The final solid was dried under
vacuum at 50 °C. Yield: 99% 1H NMR (δH; CDCl3; Me4Si): 7.00(broad, 1H), 2.80(broad, 2H), 2.64(broad, 2H), 1.65(broad, 4H), 1.53–1.22(broad, 12H), 0.88(broad, 6H). 19F NMR (δF; CDCl3; Me4Si): −139.10, −150.91, −159.94.

2.3.3. Side [P3HT0.6-(P3HT-5F)0.2-(P3HT-5F-N3)0.2] A 50 mL round bottom flask with a reflux condenser and a magnetic stirrer was degassed and filled with argon. Side [P3HT0.6-(P3HT-5F)0.4] (400 mg, 1.72 mmol) was dissolved in 35 mL THF (dry) and NaN3 (22 mg, 0.34 mmol) was added. The mixture was degassed and flushed with argon again and heated at 40 °C for 48 h. The reaction mixture was then precipitated into a mixture of deionized water:methanol (1:1). The desired azide was obtained after filtration and the solid dried under vacuum at 30 °C for 24 h. Yield: 96% 1H NMR (δH; CDCl3; Me4Si): 7.00(broad, 1H), 2.80(broad, 2H), 2.64(broad, 2H), 1.65(broad, 4H), 1.53–1.22(broad, 12H), 0.88(broad, 6H). 19F NMR (δF; CDCl3; Me4Si): −135.76, −136.36, −151.20. 15N NMR (δN; CDCl3; Me4Si): 233.84, 51.73.

The same procedure was used for the side [P3HT0.6-(P3HT-5F-N3)0.4].

2.3.4. Side [P3HT0.6-(P3HT-5F-N-C60)0.2] A 50 mL round bottom flask, equipped with a reflux condenser and a magnetic stirrer, was degassed (flamed under vacuum) and filled with argon. Side [P3HT0.6-(P3HT-5F-N-C60)0.4] (150 mg), C60 (87 mg, 0.12 mmol) and 60 mL o-DCB were added and the system was degassed and flushed with argon again. The reaction mixture was stirred at 140 °C for 72 h. After evaporation of the solvent, the solid was stirred in toluene for 72 h in order to remove the unreacted C60. The mixture was filtrated and the solid was evaluated via Thin Layer Chromatography (TLC), using toluene as the mobile phase on silica gel plates, and ATR for traces of unreacted C60 and if necessary the solid was again stirred in toluene. The final purified hybrid material was dried under vacuum at 50 °C overnight affording 100.00 mg of the hybrid C60 based polymer.

2.3.5. Side [P3HT0.6-(P3HT-5F-N-C70)0.1] A 50 mL round bottom flask, equipped with a reflux condenser and a magnetic stirrer, was degassed (flamed under vacuum) and filled with argon. Side [P3HT0.6-(P3HT-5F-N-C70)0.4] (150 mg), C70 (61 mg, 0.093 mmol) and 50 mL o-DCB were added and the system was degassed and flushed with argon again. The reaction mixture was stirred at 140 °C for 72 h. After evaporation of the solvent, the solid was stirred in toluene for 72 h in order to remove the unreacted C70. The mixture was filtrated and the solid was evaluated via TLC, using toluene as the mobile phase on silica gel plates, and ATR for traces of unreacted C70 in which case the solid was again stirred in toluene. The final purified material was dried under vacuum at 50 °C overnight affording 115.00 mg of the hybrid C70 based polymer.

3. Results and Discussion

3.1. Synthesis of Side-Br P3HTs

Typical GRIM polymerization conditions of 2,5-dibromo-3-hexyl-thiophene in freshly distilled THF using MeMgCl as the Grignard reagent and NiCl2(dppp) as catalyst were used to prepare the initial rr-P3HT [55]. The crude polymer was obtained after precipitation in methanol and was further purified via Soxhlet extractions using methanol, n-hexane and chloroform as fractionation solvents. For this work, we used only the chloroform fraction of the rr-P3HT with a corresponding Mn = 28,500, PDI = 1.2. For the side functionalization of rr-P3HT, bromination at the 4-position of the thiophene ring was performed [41,56–59] (Scheme 1). Various equivalents of the brominating reagent, N-Bromosuccinimide (NBS), in respect to the thiophene repeating units were used to achieve different bromination degrees of the polymeric backbone. The rate of bromine-functionalization is calculated from the 1H NMR of the side-[P3HT-(P3HT-Br)] (Figure 1). The appearance of a broadened bimodal peak at ~7–7.2 ppm and of two peaks at ~2.8–2.6 ppm is attributed to the successful bromination of
the thiophene ring at 4-position and also the partial disruption of the regioregularity of the P3HT’s backbone. More specifically, the insertion of the bromine atom at the 4-position of the thiophene ring causes the stepwise reduction of the aromatic hydrogens of P3HT initially located at ~7 ppm and a stepwise shift of the α-methylene hydrogen peak of the hexyl group from 2.8 to ~2.6 ppm. The initial rr-P3HT shows a peak at 2.8 ppm corresponding to the α-methylene protons of the hexyl aliphatic chains at the 3-position of the thiophene ring connected in a head-to-tail mode. After bromination at the 4-position of the thiophene ring, via electrophilic substitution affording the side-[P3HT-(P3HT-Br)] polymers (Scheme 1), a new broad proton signal at about 2.6 ppm appears attributed to the α-methylene protons of those hexyl chains attached to the partially brominated thiienyl ring. This chemical shift is down-field shifted compared to that of their initial counterparts, indicating the influence of the bromine as an electron-attracting group on the α-thienyl ring. This chemical shift is down-field shifted compared to that of their initial counterparts, attributed to the α-methylene protons of those hexyl chains attached to the partially brominated thiienyl ring. From the integral ratio of the peaks at 2.8–2.6 ppm to the one at ~7 ppm, the bromine substitution ratio in the side-[P3HT-(P3HT-Br)] polymers is calculated. In the ATR spectra (Figure 2), it is observed that as the ratio of the bromine-functionalization increases the characteristic peaks of the rr-P3HT are broadened, compared to the unfunctionalized rr-P3HT. The appearance of two new peaks at ~1600 and ~1720 cm\(^{-1}\), respectively, is attributed to the successful bromination of the thiophene ring at the 4-position [41].

![Scheme 1. Synthetic procedure of side functionalization of rr-P3HT.](image)

| x (theoretical) | Integration | H2 | H2O | x (experimental) |
|----------------|-------------|----|-----|------------------|
| 50%            | 1           | 4.122 |      | 51.07%          |
| 40%            | 1           | 3.295 |      | 40.08%          |
| 25%            | 1           | 2.320 |      | 25.16%          |
| 10%            | 1           | 2.015 |      | 10.46%          |
| 0%             | 1           | 2.045 |      | 0%              |

![Figure 1. \(^1H\) NMR spectra of the various side bromine functionalized P3HTs, in CDCl\(_3\) (the asterisks indicate solvent traces).](image)
Therefore, the side rr polymer (Scheme 2). The successful introduction of the perfluorophenyl ring at the side position of the P3HT’s polymeric backbone was proven through 19F NMR spectroscopy (Figure 3). The 19F NMR spectrum presents three peaks attributed to the three fluorine species of the perfluorinated ring. Afterwards, the perfluorophenyl groups of the P3HT0.6-(P3HT-5F)0.4 polymer were transformed into azides using sodium azide in THF [38]. Various azidation degrees were employed affording materials with 40%, 20% or 10% overall theoretical azidation degrees (Scheme 2, Table 1) in order to be able thereafter to incorporate different amounts of fullerenes and eventually produce processable hybrid materials with adequate functionalization levels and with efficient number of side-fullerene units to act as compatibilizers of the neat components’ blends, we chose the 40% side bromine-functionalized P3HT for further studies. Therefore, the side rr-[P3HT0.6-(P3HT-Br)0.4] reacted with an equivalent amount of perfluorophenyl boronic acid under Suzuki cross coupling conditions to form the respective side [P3HT0.6-(P3HT-5F)0.4] polymer (Scheme 2). The successful introduction of the perfluorophenyl ring at the side position of the P3HT’s polymeric backbone was proven through 19F NMR spectroscopy (Figure 3). The 19F NMR spectrum presents three peaks attributed to the three fluorine species of the perfluorinated ring. Afterwards, the perfluorophenyl groups of the P3HT0.6-(P3HT-5F)0.4 polymer were transformed into azides using sodium azide in THF [38]. Various azidation degrees were employed affording materials with 40%, 20% or 10% overall theoretical azidation degrees (Scheme 2, Table 1) in order to be able thereafter to incorporate different amounts of fullerenes and eventually produce processable hybrid polymers with C60 or C70. The successful azidation was confirmed using ATR and 15N NMR spectroscopies. The ATR spectrum of the 20% side perfluorophenyl azide showed a low intensity peak at 2064 cm−1 which demonstrates the azide formation (Figure 4), while for the 10% azide functionalized polymer the azide peak was not clearly observed due to the low ratio of the azide functionality over the total polymer. On the other hand, 15N NMR spectroscopy, as shown in Figure 5, revealed two clear peaks owing to the two nitrogen species of the azide functionality even for the 10% azide functionalized polymer.  

As it is known from previous literature findings on the side functionalization of rr-P3HT [41,56–59], high levels of functionalization at the 4-position of the thiophene ring affect the planarity along the polymer’s backbone and also disrupt the effective conjugation length of P3HT. Additionally, high functionalization degrees with fullerene species along the rr-P3HT backbone, which is our herein ultimate target, were inevitably going to provide hybrid materials with poor solubility in common organic solvents. Thus, and in order to prepare processable hybrid materials, but with adequate functionalization levels and with efficient number of side-fullerene units to act as compatibilizers of the neat components’ blends, we chose the 40% side bromine-functionalized P3HT for further studies.  

![Figure 2. ATR spectra of an unfunctionalized P3HT and various ratios of side bromine-functionalized P3HTs.](image)

Table 1. Azidation ratios of [P3HT0.6-(P3HT-5F)0.4] and the respective abbreviations of the functionalized polymers obtained.

| Azide Functionalization Degree of (P3HT-5F-N3)x-(P3HT-5F)ax-(P3HT)1-x | Side azide |
|---|---|
| 1 | [P3HT0.6-(P3HT-5F-N3)0.4] |
| 0.5 | [P3HT0.6-(P3HT-5F)0.2-(P3HT-5F-N3)0.2] |
| 0.25 | [P3HT0.6-(P3HT-5F)0.3-(P3HT-5F-N3)0.1] |
Scheme 2. Synthetic procedure of side perfluorophenyl P3HT and their azide transformation; $a$ indicates the azide functionalization degree.

Figure 3. $^{19}$F NMR spectra (in CDCl$_3$) of the P3HT$_{0.6}$-(P3HT-5F)$_{0.4}$.

Figure 4. ATR spectra of the [P3HT$_{0.6}$-(P3HT-5F)$_{0.4}$] and azide [P3HT$_{0.6}$-(P3HT-5F)$_{0.2}$-(P3HT-5F-N$_3$)$_{0.2}$] (the asterisk indicates the azide functionality).

Figure 5. $^{15}$N NMR spectra of the side [P3HT$_{0.6}$-(P3HT-5F)$_{0.3}$-P3HT-5F-N$_3$)$_{0.1}$], in CDCl$_3$. 
3.2. Synthesis of Side-P3HT-(P3HT-5F) Hybrids

The side P3HT-perfluorophenyl azides have been used to prepare side hybrid semiconducting polymers with either C\textit{60} or C\textit{70} fullerenes. When the \textit{side} [P3HT\textsubscript{0.6}-(P3HT-5F-N\textsubscript{3})\textsubscript{0.4}] reacted with an equivalent amount of C\textit{60} to the azide content, the final hybrid polymer was insoluble in common organic solvents. In order to avoid such insolubility problems, the \textit{side} [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.2}-(P3HT-5F-N\textsubscript{3})\textsubscript{0.2}] reacted with C\textit{60} fullerene while the \textit{side} [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N\textsubscript{3})\textsubscript{0.1}] reacted with C\textit{70} providing the desired hybrid semiconducting polymers (Scheme 3) and leading to processable hybrid materials. After the [3+2] cycloaddition reaction, the \textit{side} P3HT-(P3HT-5F) hybrids with fullerenes were thoroughly washed several times with toluene to remove any unreacted fullerene species. A thorough ATR examination was performed for both hybrid materials after each subsequent washing step. When no further reduction of the fullerene peaks was noticed, we concluded that all unreacted traces of C\textit{60} or C\textit{70} were successfully removed (Figure 6).

![Scheme 3. Synthetic procedure of side-[P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] hybrids with C\textit{60} or C\textit{70} fullerenes.](image)

![Figure 6. ATR spectra of: (a) C\textit{60} and (b) C\textit{70} based side [(P3HT\textsubscript{0.6}-P3HT-5F)\textsubscript{0.4}] hybrids, in comparison with the initial P3HT-(P3HT-5F) and the plain fullerenes, respectively.](image)
and a 62% residue for the [P3HT\textsubscript{0.6}-\{(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}\}] hybrid, respectively. As expected, the hybrid polymer presented a higher residue due to the incorporation of the fullerene species.

3.3. Optical Properties

The optical properties of the synthesized hybrid materials compared to the initial side [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] were investigated both in o-DCB solutions and in film form. The UV-Vis spectra of the initial side-[P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] both in o-DCB solution (Figure 8a) and in film form (Figure 8a) presented a blue shift compared to the unfunctionalized rr-P3HT. This phenomenon is attributed to the perturbation of the regioregularity of the P3HT backbone after the functionalization at the 4-position of the thiophene unit. The hybrid material [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.2}-(P3HT-5F-N-C\textsubscript{60})\textsubscript{0.2}] presented an even greater blue-shift compared to its functionalized P3HT precursor, while the characteristic absorbance bands of the fullerene C\textsubscript{60} were observed in solution as a shoulder at \~330 nm whereas, in film form, the absorption peak up to \~300 nm peak is attributed to the fullerene part of the hybrid. On the other hand, the hybrid material [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}] clearly presented the characteristic absorbance bands of C\textsubscript{70} fullerene both in solutions and in film form in the region 300–400 nm.

![Figure 7. TGA thermograms of rr-P3HT, [P3HT\textsubscript{0.6}-(P3HT-Br)\textsubscript{0.4}], [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}], [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}], and C\textsubscript{70} fullerene.](image)

![Figure 8. Normalized Absorption spectra in: (a) o-DCB solutions; and (b) film form of unfunctionalized P3HT, side functionalized [P3HT\textsubscript{0.6}-(P3HT-Br)\textsubscript{0.4}], [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}], [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.2}-(P3HT-5F-N-C\textsubscript{60})\textsubscript{0.2}], and [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}], respectively. The inset in (a) shows the Normalized Absorption spectra of C\textsubscript{60} and C\textsubscript{70} in o-DCB solutions.](image)
The PL spectra of side-[P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] and its respective hybrids with C\textsubscript{60} and C\textsubscript{70} both in o-DCB solutions and in film form are shown in Figure 9. The o-DCB solutions of the hybrid materials, upon excitation at 470 nm, presented blue shifted spectra and broader photoluminescence curves in the region 500–650 nm, compared to the un-functionalized P3HT (Figure 9a). In film form, after excitation at the UV-Vis films maxima, the PL spectra followed the trend of the respective solutions also showing a blue shift compared to the neat rr-P3HT. A small quenching of the hybrids’ photoluminescence intensity in film form was observed, [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.2}-(P3HT-5F-N-C\textsubscript{60})\textsubscript{0.2}] green-colored line and [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}] blue-colored line of Figure 9b, that is attributed to the electron accepting fullerene parts interacting electronically with the electron donating P3HT polymeric backbone and the perturbation of the P3HT’s backbone due to the side functionalization.

3.4. Morphology Characterization

Thin film morphology of side perfluorophenyl functionalized [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] and its respective hybrids with C\textsubscript{60} and C\textsubscript{70} was investigated using transmission electron spectroscopy (TEM) (Figure 10). The initial side [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}] formed a uniform thin film in the nanometer scale. The corresponding C\textsubscript{60} based hybrid, [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.2}-(P3HT-5F-N-C\textsubscript{60})\textsubscript{0.2}], formed films with large domains compared to the initial perfluorophenyl modified P3HT, with darker and brighter regions. In contrast, the C\textsubscript{70} based hybrid polymer, [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.3}-(P3HT-5F-N-C\textsubscript{70})\textsubscript{0.1}], presented a uniform nanophase separated thin film formation with well resolved small domains in analogy to its non hybrid initial polymer, [P3HT\textsubscript{0.6}-(P3HT-5F)\textsubscript{0.4}].
Moreover, a typical P3HT:PC$_{70}$BM blend of 1:1 $w/w$ ratio was employed to prepare two new blends with the [P3HT$_{0.6}$-(P3HT-5F)$_{0.3}$-(P3HT-5F-N-C$_{70}$)$_{0.1}$] hybrid polymer. The hybrid material was added at a 20 wt % and at a 50 wt % percent, respectively, over the total weight of the P3HT:PC$_{70}$BM initial blend. The effect of the additive on the morphology of the initial blend was investigated via TEM microscopy. Figure 11 presents the respective TEM images in 100 nm and in 20 nm scale, of the initial blend and of the blends containing the hybrid polymer without any thermal or other solvent vapor treatment. The P3HT:PC$_{70}$BM blend presented uniform thin film with nanophase separation. The morphology of the blend containing 20% of the side C$_{70}$ based hybrid polymer did not present any differentiations, even at the 20 nm scale, showing the probable stabilization of the initial blends morphology. On the other hand, the blend with the 50% of the side C$_{70}$ hybrid polymer formed more heterogeneous films in which darker domains are present.

![Figure 11. TEM images of a P3HT:PC$_{70}$BM 1:1 $w/w$ blend and of P3HT:PC$_{70}$BM: [P3HT$_{0.6}$-(P3HT-5F)$_{0.3}$-(P3HT-5F-N-C$_{70}$)$_{0.1}$] blends with a 20 wt % and a 50 wt % of the [P3HT$_{0.6}$-(P3HT-5F)$_{0.3}$-(P3HT-5F-N-C$_{70}$)$_{0.1}$] hybrid, respectively. For the top images, the scale bars correspond to 100 nm while, for the bottom images, they correspond to 20 nm.]

4. Conclusions

In this work, modification in various ratios of the $rr$-P3HT backbone in order to introduce the perfluorophenyl functionality at the 4-position of the thiophene unit has been efficiently performed. Thereafter, and based on our previous reported methodology, the perfluorophenyl moieties transformed into azides, from which hybrid materials with fullerenes, C$_{60}$ or C$_{70}$, have been synthesized aiming to develop efficient compatibilizers and stabilizers of P3HT:fullerene active layers typically employed in large area flexible OPVs. The side-functionalized P3HTs and their respective hybrid polymers have been characterized in respect to their optical and morphological properties. In solutions and in film form, all the side-functionalized materials presented blue shifted absorbance curves due to the perturbation of regioregularity of polythiophene. Regarding their photoluminescence in film form, the hybrid materials presented moderate photoluminescence quenching. The C$_{70}$ based hybrid polymer formed uniform nanophase separated thin films while the C$_{60}$ based hybrid formed more heterogeneous thin films exhibiting larger domains. In order to investigate their applicability as potential compatibilizers of polythiophene:fullerene BHJ active layers, selected hybrids were evaluated for their influence on the morphology of a typical P3HT:PC$_{70}$BM active layer blend. More specifically, when the percentage of the hybrid was low compared to the total blend weight ratio, the morphology...
was not affected. However, when the hybrid polymer was added at a higher percentage, the final blend presented irregularities and large domains were created throughout the blend.

Based on these findings it can be concluded that the particular “perfluorophenyl”-“azide”-“carbon nanostructure hybrid” methodology is conveniently applicable for the development of side hybrid semiconducting polymer electron donors allowing the insertion of fullerene species in the desired percentage. Such hybrid polymer–fullerene materials can be employed as additives of typical polymer-donor:fullerene-acceptor active layer blends for OPVs.

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References
1. Yu, J.; Zheng, Y.; Huang, J. Towards high performance organic photovoltaic cells: A review of recent development in organic photovoltaics. Polymers 2014, 6, 2473–2509. [CrossRef]
2. Mazzio, K.A.; Luscombe, C.K. The future of organic photovoltaics. Chem. Soc. Rev. 2015, 44, 78–90. [CrossRef] [PubMed]
3. Lu, L.; Zheng, T.; Wu, Q.; Schneider, A.M.; Zhao, D.; Yu, L. Recent advances in bulk heterojunction polymer solar cells. Chem. Rev. 2015, 115, 12666–12731. [CrossRef] [PubMed]
4. Ma, W.; Yang, C.; Gong, X.; Lee, K.; Heeger, A.J. Thermally stable, efficient polymer solar cells with nanoscale control of the interpenetrating network morphology. Adv. Funct. Mater. 2005, 15, 1617–1622. [CrossRef]
5. Choulis, S.A.; Kim, Y.; Nelson, J.; Bradley, D.D.C.; Giles, M.; Shkunov, M.; McCulloch, I. High ambipolar and balanced carrier mobility in regioregular poly(3-hexylthiophene). Appl. Phys. Lett. 2004, 85, 3890–3892. [CrossRef]
6. Schafferhans, J.; Baumann, A.; Wagenpfahl, A.; Deibel, C.; Dyakonov, V. Oxygen doping of P3HT:PCBM blends: Influence on trap states, charge carrier mobility and solar cell performance. Org. Electron. 2010, 11, 1693–1700. [CrossRef]
7. Dang, M.T.; Wantz, G.; Bejbouji, H.; Urien, M.; Daulet, O.J.; Vignau, L.; Hirsch, L. Polymeric solar cells based on P3HT:PCBM: Role of the casting solvent. Sol. Energy Mater. Sol. Cells 2011, 95, 3408–3418. [CrossRef]
8. Baek, W.-H.; Yang, H.; Yoon, T.-S.; Kang, C.J.; Lee, H.H.; Kim, Y.-S. Effect of P3HT:PCBM concentration in solvent on performances of organic solar cells. Sol. Energy Mater. Sol. Cells 2009, 93, 1263–1267. [CrossRef]
9. Carlé, J.E.; Krebs, F.C. Technological status of organic photovoltaics (OPV). Sol. Energy Mater. Sol. Cells 2013, 119, 309–310. [CrossRef]
10. Kapnopoulou, C.; Mekeridis, E.D.; Tzounis, L.; Polyzoidis, C.; Zachariadis, A.; Tsimikli, S.; Gravalidis, C.; Laskarakis, A.; Vouroutzis, N.; Logothetidis, S. Fully gravure printed organic photovoltaic modules: A straightforward process with a high potential for large scale production. Sol. Energy Mater. Sol. Cells 2016, 144, 724–731. [CrossRef]
11. Kallitsis, J.K.; Kourkouli, S.; Andreopoulou, A.K. Organic Polymeric Semiconductor Materials for Applications in Photovoltaic Cells Part 1: Properties and Materials, Handbook on Flexible Organic Electronics: Materials, Manufacturing and Applications; Logothetidis, S., Ed.; Woodhead Publishing Limited: Cambridge, UK, 2015; pp. 85–114.
12. Scharber, M.C.; Mühlbacher, D.; Koppe, M.; Denk, P.; Waldauf, C.; Heeger, A.J.; Brabec, C.J. design rules for donors in bulk-heterojunction solar cells—Towards 10% energy-conversion efficiency. Adv. Mater. 2006, 18, 789–794. [CrossRef]
13. Liu, X.; Chen, H.; Tan, S. Overview of high-efficiency organic photovoltaic materials and devices. Renew. Sustain. Energy Rev. 2015, 52, 1527–1538. [CrossRef]
14. Gao, M.; Subbiah, J.; Geraghty, P.B.; Chen, M.; Purushothaman, B.; Chen, X.; Qin, T.; Vak, D.; Scholes, F.H.; Watkins, S.E.; et al. Development of a high-performance donor–acceptor conjugated polymer: Synergy in materials and device optimization. *Chem. Mater.* 2016, 28, 3481–3487. [CrossRef]

15. Chen, J.; Cao, Y. Development of novel conjugated donor polymers for high efficiency bulk-heterojunction photovoltaic devices. *Acc. Chem. Res.* 2009, 42, 1709–1718. [CrossRef] [PubMed]

16. He, Y.; Li, Y. Fullerene derivative acceptors for high performance polymer solar cells. *Phys. Chem. Chem. Phys.* 2011, 13, 1970–1983. [CrossRef] [PubMed]

17. He, Y.; Chen, H.-Y.; Hou, J.; Li, Y. Indene-C60 bisadduct: A new acceptor for high-performance polymer solar cells. *J. Am. Chem. Soc.* 2010, 132, 1377–1382. [CrossRef] [PubMed]

18. He, Y.; Zhao, G.; Peng, B.; Li, Y. High-yield synthesis and electrochemical and photovoltaic properties of indene-C70 bisadduct. *Adv. Funct. Mater.* 2010, 20, 3383–3389. [CrossRef] [PubMed]

19. Facchetti, A. Polymer donor–polymer acceptor (all-polymer) solar cells. *Mater. Today* 2013, 16, 123–132. [CrossRef]

20. Bin, H.; Zhang, Z.-G.; Gao, L.; Chen, S.; Zhong, L.; Xue, L.; Yang, C.; Li, Y. Non-fullerene polymer solar cells based on alkylthio and fluorine substituted 2D-conjugated polymers reach 9.5% efficiency. *J. Am. Chem. Soc.* 2016, 138, 4657–4664. [CrossRef] [PubMed]

21. Kim, Y.; Lim, E. Development of polymer acceptors for organic photovoltaic cells. *Polymers* 2014, 6, 382–407. [CrossRef]

22. Brady, M.A.; Su, G.M.; Chabinyc, M.L. Recent progress in the morphology of bulk heterojunction photovoltaics. *Soft Matter* 2011, 7, 11065–11077. [CrossRef]

23. Ruderer, M.A.; Muller-Buschbaum, P. Morphology of polymer-based bulk heterojunction films for organic photovoltaics. *Soft Matter* 2011, 7, 5482–5493. [CrossRef]

24. Zhao, Y.; Xie, Z.; Qu, Y.; Geng, Y.; Wang, L. Solvent-vapor treatment induced performance enhancement of poly(3-hexylthiophene): Methanofullerene bulk-heterojunction photovoltaic cells. *Appl. Phys. Lett.* 2007, 90, 043504. [CrossRef]

25. Moulé, A.J.; Meerholz, K. Controlling morphology in polymer–fullerene mixtures. *Adv. Mater.* 2008, 20, 240–245. [CrossRef]

26. Yao, Y.; Hou, J.; Xu, Z.; Li, G.; Yang, Y. Effects of solvent mixtures on the nanoscale phase separation in polymer solar cells. *Adv. Funct. Mater.* 2008, 18, 1783–1789. [CrossRef]

27. Angiolini, L.; Cocchi, V.; Lanzi, M.; Salatelli, E.; Tonelli, D.; Vlamidis, Y. Novel regioregular polythiophenes containing side-chain porphyrin groups for polymeric photovoltaic cells. *Mat. Chem. Phys.* 2014, 146, 464–471. [CrossRef]

28. Bertho, S.; Campo, B.; Piersimoni, F.; Spoltore, D.; D’Haen, J.; Lutsen, L.; Maes, W.; Vanderzande, D.; Manca, J. Improved thermal stability of bulk heterojunctions based on side-chain functionalized poly(3-alkylthiophene) copolymers and PCBM. *Sol. Energy Mater. Sol. Cells* 2013, 110, 69–76. [CrossRef]

29. Jaymand, M.; Hatamzadeh, M.; Omid, Y. Modification of polythiophene by the incorporation of processable polymeric chains: Recent progress in synthesis and applications. *Prog. Polyom. Sci.* 2015, 47, 26–69. [CrossRef]

30. Lobez, J.M.; Andrew, T.L.; Bulović, V.; Swager, T.M. Improving the performance of P3HT–fullerene solar cells with side chain functionalized poly(thiophene) additives: A new paradigm for polymer design. *ACS Nano* 2012, 6, 3044–3056. [CrossRef] [PubMed]

31. Wang, H.-J.; Chen, C.-P.; Jeng, R.-J. Polythiophenes comprising conjugated pendants for polymer solar cells: A review. *Materials* 2014, 7, 2411–2439. [CrossRef]

32. Valderrama-García, B.X.; Rodríguez-Alba, E.; Morales-Espinoza, E.G.; Moineau Chane-Ching, K.; Rivera, E. Synthesis and characterization of novel polythiophenes containing pyrene chromophores: Thermal, optical and electrochemical properties. *Molecules* 2016, 21, 172. [CrossRef] [PubMed]

33. Liu, J.; Zhu, X.; Li, J.; Shen, J.; Tu, G. Enhancing the thermal stability of the bulk-heterojunction photovoltaics based on P3HT/PCBM by incorporating diblock amphipathic P3HT–PEO at D/A interface. *RSC Adv.* 2016, 6, 61934–61943. [CrossRef]

34. Rumer, J.W.; McCulloch, I. Organic photovoltaics: Crosslinking for optimal morphology and stability. *Mater. Today* 2015, 18, 425–435. [CrossRef]

35. Wantz, G.; Derue, L.; Daulet, O.; Rivatou, A.; Hudhomme, P.; Dagron-Lartigau, C. Stabilizing polymer-based bulk heterojunction solar cells via crosslinking. *Polym. Int.* 2014, 63, 1346–1361. [CrossRef]
36. Li, F.; Yager, K.G.; Dawson, N.M.; Jiang, Y.-B.; Malloy, K.J.; Qin, Y. Nano-structuring polymer/fullerene composites through the interplay of conjugated polymer crystallization, block copolymer self-assembly and complementary hydrogen bonding interactions. Polym. Chem. 2015, 6, 721–731. [CrossRef]

37. Huang, P.-T.; Chen, Y.-H.; Lin, B.-Y.; Chuang, W.-P. Homogenized poly(3-hexylthiophene)/methanofullerene film by addition of end-functionalized compatibilizer and its application to polymer solar cells. Int. J. Photoenergy 2015, 2015, 762532. [CrossRef]

38. Kim, H.J.; Han, A.-R.; Cho, C.-H.; Kang, H.; Cho, H.-H.; Lee, M.Y.; Fréchet, J.M.J.; Oh, J.H.; Kim, B.J. Solvent-resistant organic transistors and thermally stable organic photovoltaics based on cross-linkable conjugated polymers. Chem. Mater. 2012, 24, 215–221. [CrossRef]

39. Yuan, K.; Chen, L.; Chen, Y. Nanostructuring compatibilizers of block copolymers for organic photovoltaics. Polym. Int. 2014, 63, 593–606. [CrossRef]

40. Kallitsis, J.K.; Anastasopoulos, C.; Andreopoulou, A.K. Functional semiconductors targeting copolymer architectures and hybrid nanostructures. MRS Commun. 2015, 5, 365–382. [CrossRef]

41. Li, M.; Xu, P.; Yang, J.; Yang, S. Donor-p-acceptor double-cable polythiophenes bearing fullerene pendant with tunable donor/acceptor ratio: A facile postpolymerization. J. Mater. Chem. 2010, 20, 3953–3960. [CrossRef]

42. Gholamkhass, B.; Peckham, T.J.; Holdcroft, S. Poly(3-hexylthiophene) bearing pendant fullerenes: Aggregation vs. self-organization. Polym. Chem. 2010, 1, 708–719. [CrossRef]

43. Lanzi, M.; Salatelli, E.; Benelli, T.; Caretti, D.; Giorgini, L.; Di-Nicola, F.P. A regioregular polythiophene-fullerene with tunable donor/acceptor ratio: A facile postpolymerization. J. Mater. Chem. Sci. 2015, 132. [CrossRef]

44. Hufnagel, M.; Fischer, M.; Thurm-Albrecht, T.; Thelakkat, M. Influence of fullerene grafting density on structure, dynamics, and charge transport in P3HT-b-PPC61BM block copolymers. Macromolecules 2016, 49, 1637–1647. [CrossRef]

45. Raïssi, M.; Erothu, H.; Ibarboure, E.; Cramail, H.; Vignau, L.; Cloutet, E.; Hiorns, R.C. Fullerene-capped copolymers for bulk heterojunctions: Device stability and efficiency improvements. J. Mater. Chem. A 2015, 3, 18207–18221. [CrossRef]

46. Kourkouli, S.N.; Siokou, A.; Stefopoulos, A.A.; Ravani, F.; Plocke, T.; Müller, M.; Maultzsch, J.; Thomsen, C.; Papagelis, K.; Kallitsis, J.K. Electronic properties of semiconducting polymer-functionalized single wall carbon nanotubes. Macromolecules 2013, 46, 2590–2598. [CrossRef]

47. Stefopoulos, A.A.; Chochos, C.L.; Prato, M.; Pistolis, G.; Papagelis, K.; Petraki, F.; Kennou, S.; Kallitsis, J.K. Novel hybrid materials consisting of regioregular poly(3-octylthiophene) covalently attached to single wall carbon nanotubes. Chem. Eur. J. 2008, 14, 8715–8724. [CrossRef] [PubMed]

48. Stefopoulos, A.A.; Kourkouli, S.N.; Economopoulos, S.; Ravani, F.; Andreopoulou, A.; Papagelis, K.; Siokou, A.; Kallitsis, J.K. Polymer and hybrid electron accepting materials based on a semiconducting perfluorophenylquinoline. Macromolecules 2010, 43, 4827–4828. [CrossRef]

49. Kakogianni, S.; Kourkouli, S.N.; Andreopoulou, A.K.; Kallitsis, J.K. A versatile approach for creating hybrid semiconducting polymer–fullerene architectures for organic electronics. J. Mater. Chem. A 2014, 2, 8110–8117. [CrossRef]

50. Sygellou, L.; Kakogianni, S.; Andreopoulou, A.K.; Theodossiou, K.; Leftheriotis, G.; Kallitsis, J.K.; Siokou, A. Evaluation of the electronic properties of perfluorophenyl functionalized quinolines and their hybrids with carbon nanostructures. Phys. Chem. Chem. Phys. 2016, 18, 4154–4165. [CrossRef] [PubMed]

51. Raja, R.; Liu, W.-S.; Hsiow, C.-Y.; Rwei, S.-P.; Chiu, W.-Y.; Wang, L. Terthiophene-C60 dyad as donor/acceptor compatibilizer for developing highly stable P3HT/PC61BM bulk heterojunction solar cells. J. Mater. Chem. A 2015, 3, 14401–14408. [CrossRef]

52. Perrin, L.; Legros, M.; Mercier, R. Design of a series of polythiophenes containing C60 groups: Synthesis and optical and electrochemical properties. Macromolecules 2015, 48, 323–336. [CrossRef]

53. Yameen, B.; Puerckhauer, T.; Ludwig, J.; Ahmed, I.; Altintas, O.; Fruk, L.; Colmann, A.; Barner-Kowollik, C. π-conjugated polymer–fullerene covalent hybrids via ambient conditions diels–alder ligation. Small 2014, 10, 3091–3098. [CrossRef] [PubMed]

54. Kakogianni, S.; Lebedeva, M.A.; Paloumbis, G.; Andreopoulou, A.K.; Porfyrikis, K.; Kallitsis, J.K. Semiconducting end-perfluorinated P3HT-fullerene hybrids as potential additives for P3HT/IC70BA blends. RSC Adv. 2016, 6, 98306–98316. [CrossRef]
55. Loewe, R.S.; Ewbank, P.C.; Liu, J.; Zhai, L.; McCullough, R.D. Regioregular, head-to-tail coupled poly(3-alkylthiophenes) made easy by the GRIM method: Investigation of the reaction and the origin of regioselectivity. *Macromolecules* 2001, 34, 4324–4333. [CrossRef]

56. Chochos, C.L.; Economopoulos, S.P.; Deimede, V.; Gregoriou, V.G.; Lloyd, M.T.; Malliaras, G.G.; Kallitsis, J.K. Synthesis of a soluble n-type cyano substituted polythiophene derivative: A potential electron acceptor in polymeric solar cells. *J. Phys. Chem. C* 2007, 111, 10732–10740. [CrossRef]

57. Economopoulos, S.P.; Chochos, C.L.; Gregoriou, V.G.; Kallitsis, J.K.; Barrau, S.; Hadziioannou, G. Novel brush-type copolymers bearing thiophene backbone and side chain quinoline blocks. Synthesis and their use as a compatibilizer in thiophene-quinoline polymer blends. *Macromolecules* 2007, 40, 921–927. [CrossRef]

58. Li, Y.N.; Vamvounis, G.; Holdcroft, S. Facile functionalization of poly(3-alkylthiophene)s via electrophilic substitution. *Macromolecules* 2001, 34, 141–143. [CrossRef]

59. Li, Y.N.; Vamvounis, G.; Yu, J.; Holdcroft, S. A novel and versatile methodology for functionalization of conjugated polymers. Transformation of poly(3-bromo-4-hexylthiophene) via palladium-catalyzed coupling chemistry. *Macromolecules* 2001, 34, 3130–3132. [CrossRef]

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