

Supporting Information

Poly(3-hexylthiophene)-block-poly(tetrabutylammonium-4-styrenesulfonate) Block Copolymer Micelles for the Synthesis of Polymer Semiconductor Nanocomposites

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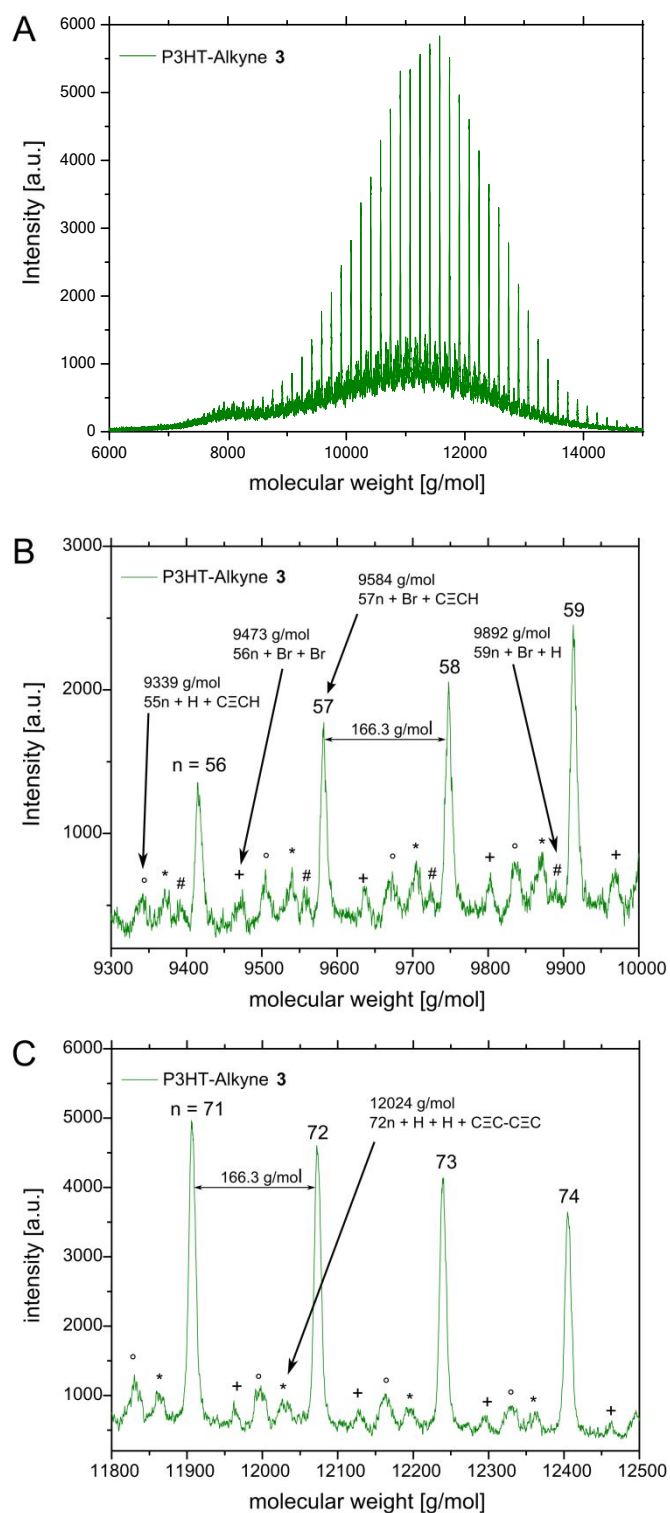


Figure S1. MALDI-ToF of P3HT-Alkyne with the whole molecular weight distribution (A) and detailed analysis of signals for polymer chains with 56-59 repeating units (B) and 71-74 repeating units (C).

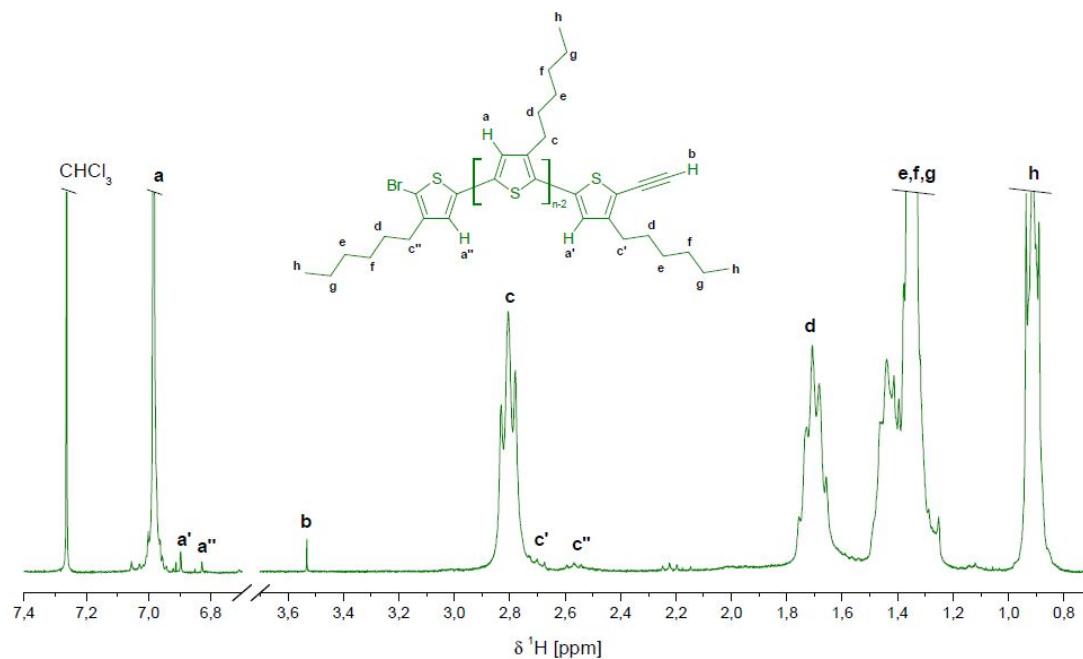


Figure S2. ^1H -NMR of P3HT-Alkyne, the signal of the alkyne proton b can be seen at $\delta = 3.51$ ppm.

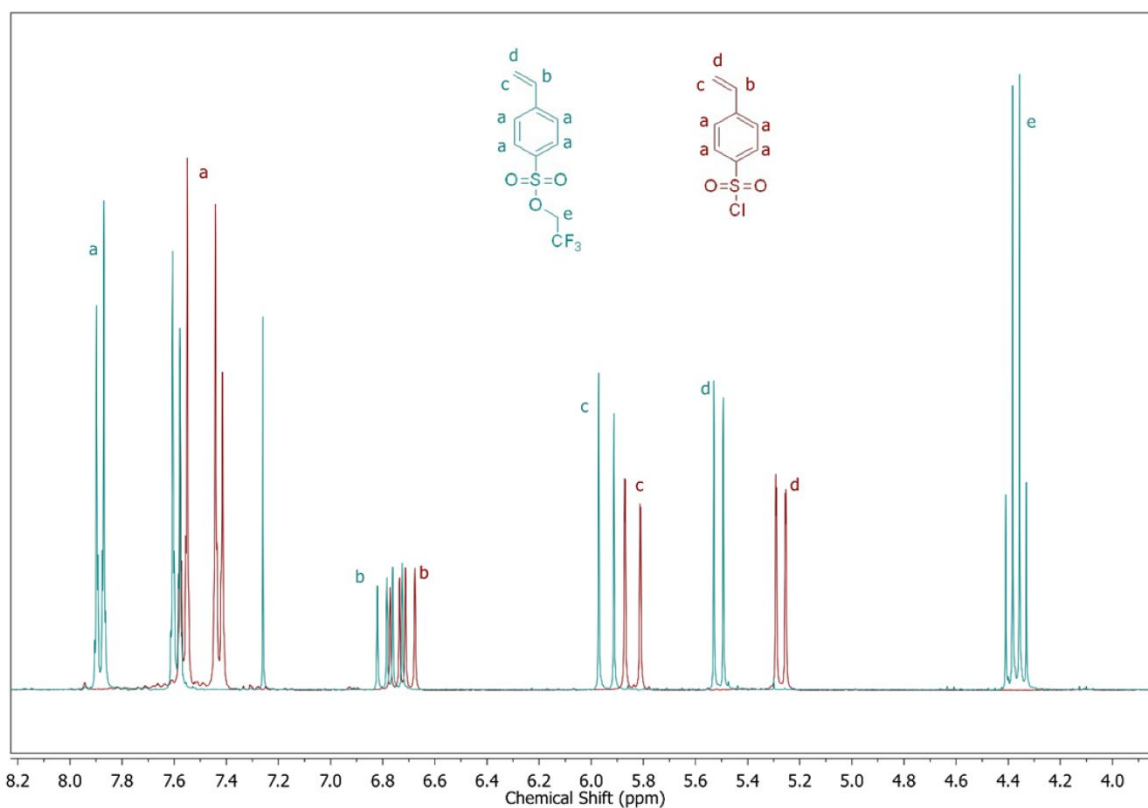


Figure S3. ^1H -NMR of 4-vinylbenzene-1-sulfonyl chloride (red) and 2,2,2-trifluoroethyl-4-vinylbenzenesulfonate (green). The overlay of the two spectra shows the complete conversion during the synthesis.

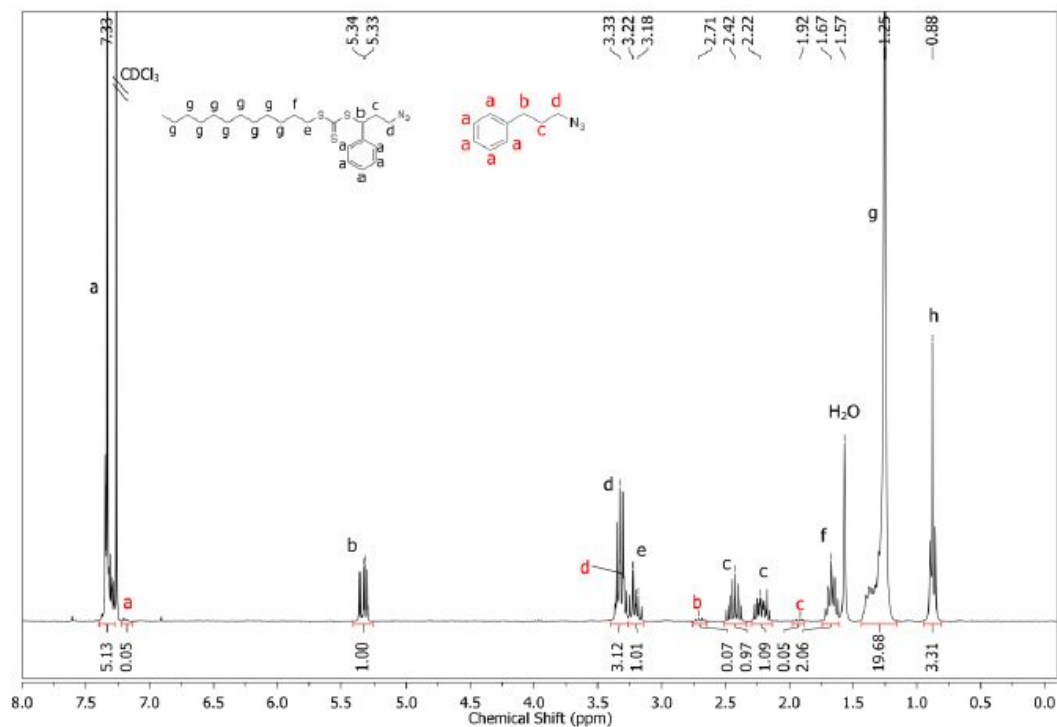


Figure S4. ^1H -NMR of 3-azido-1-phenylpropyl dodecyl trithiocarbonate (black) where small impurities (>5 mol-%) of the educt 3-azidopropylbenzene (red) are still present after purification by column chromatography.

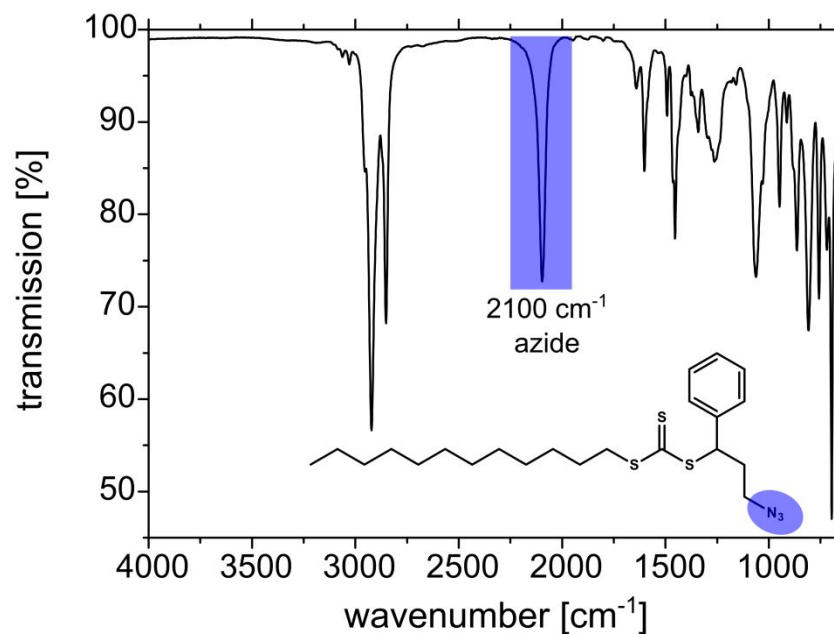


Figure S5. FT-IR spectrum of 3-azido-1-phenylpropyl dodecyl trithiocarbonate where the intense azide band is clearly visible at 2100 cm^{-1} and proves the successful synthesis of the clickable RAFT agent.

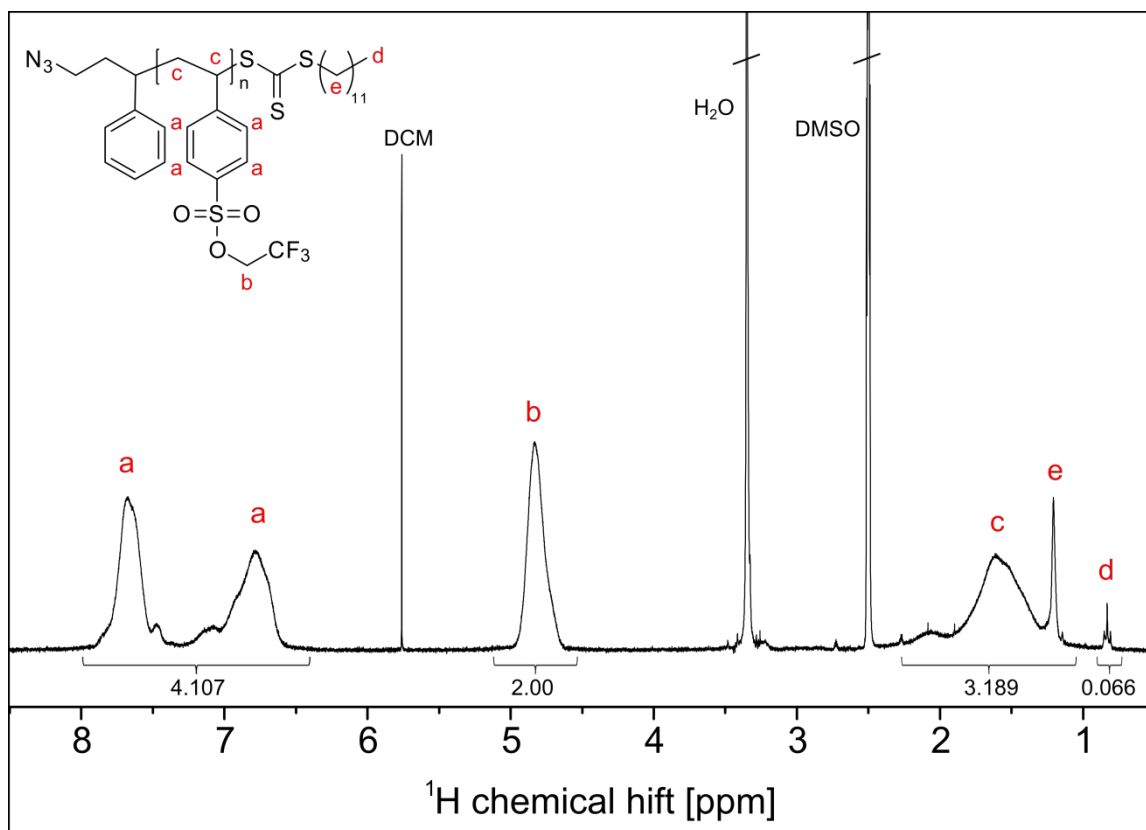


Figure S6. ^1H -NMR of $\text{P}(\text{SS}-\text{CH}_2-\text{CF}_3)-\text{N}_3$ measured in $d\text{-DMSO}$. The proton signals next the azide end group cannot be seen, because they are of small intensity and hidden under the backbone signals.

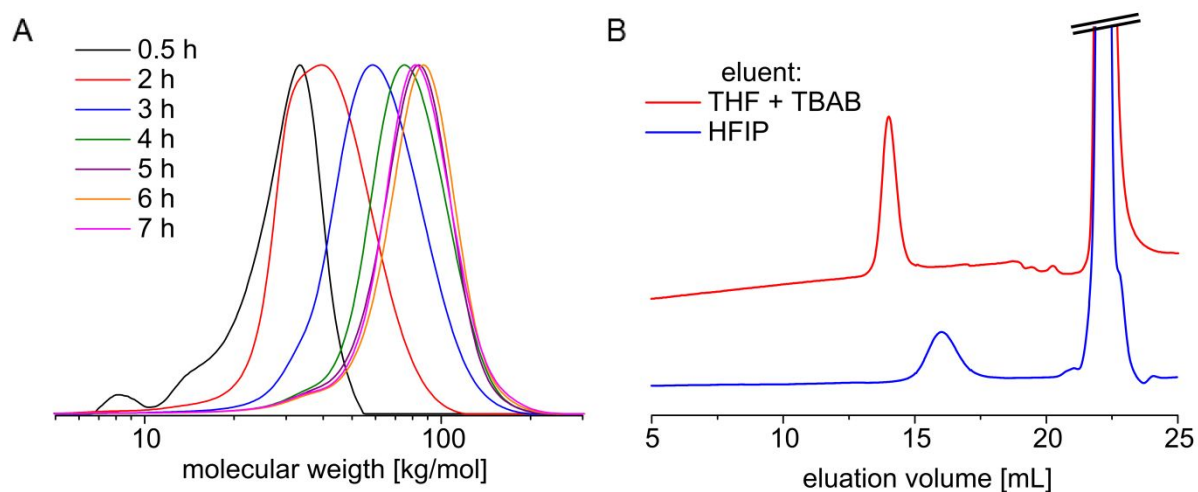


Figure S7. (A) Molecular weight distributions of $\text{P}(\text{SS}-\text{CH}_2-\text{CF}_3)-\text{N}_3$ during polymerization and (B) comparison of elugrams of $\text{P}(\text{SS}-\text{CH}_2-\text{CF}_3)-\text{N}_3$ with the solvents HFIP (toluene as internal standard) and THF + TBAB (o-dichlorobenzene as internal standard).

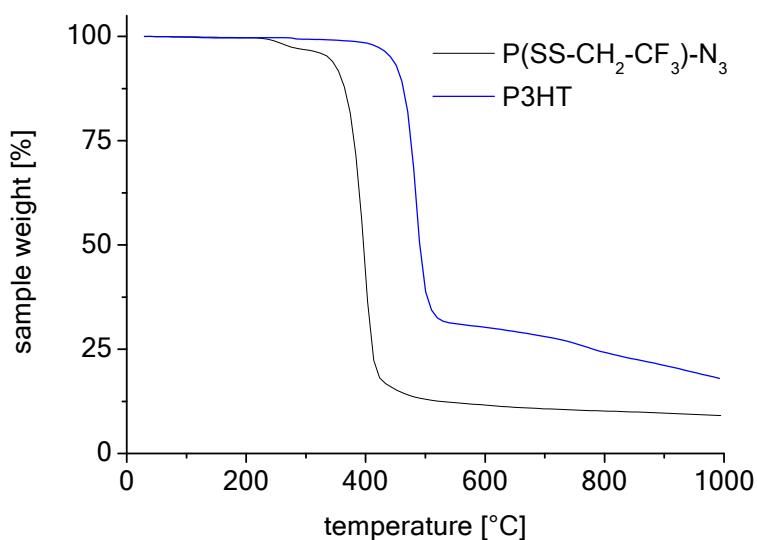


Figure S8. TGA analysis of P3HT-Alkyne and P(SS-CH₂-CF₃)-N₃ measured under N₂ atmosphere.

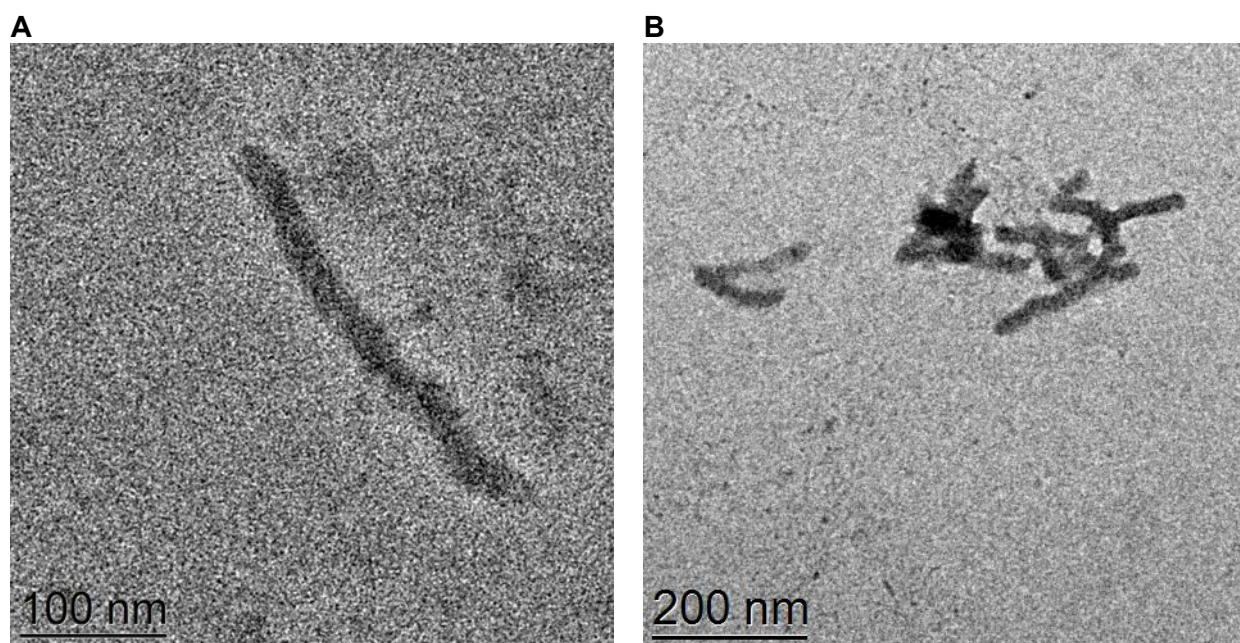


Figure S9. TEM images of P3HT-b-PSS/TBA⁺ micelles. The micelles have a wormlike structure (A) and branched shapes can be a result of agglomeration during drying (B). Samples were prepared by putting the micelle solution on the TEM-grid and remove the solvent without staining.

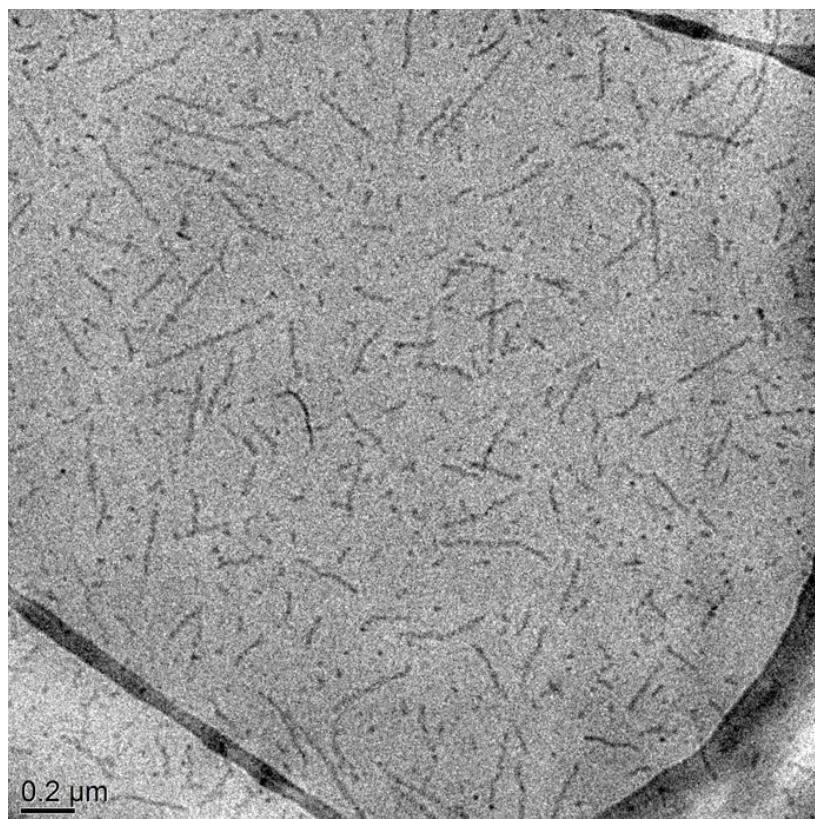


Figure S10. cryo-TEM images of P3HT-b-PSS/TBA⁺ micelles without staining (overview).

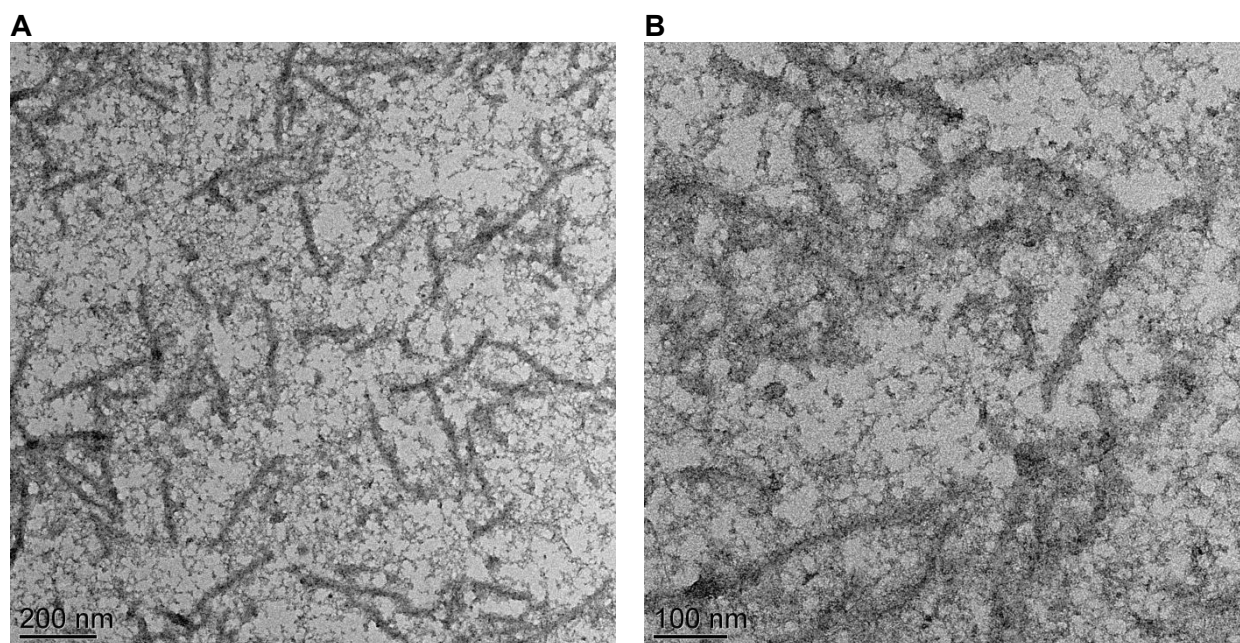


Figure S11. TEM images of dried P3HT-b-PSS/TBA⁺ + TiO₂ nanocomposites without staining.

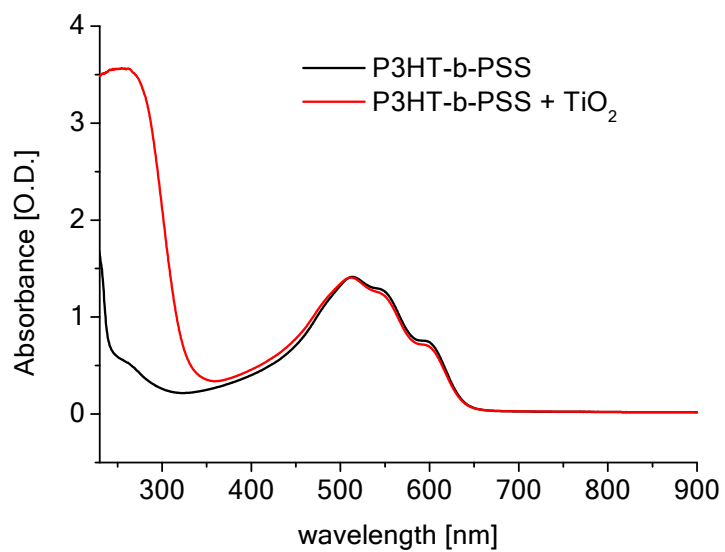


Figure S12. UV-vis spectra of P3HT-b-PSS/TBA⁺ micelles in water and P3HT-b-PSS/TBA⁺ + TiO₂ in water.