Supporting Information

Synthesis of Conjugated Polymers *via* Exclusive Direct-Arylation Coupling Reaction: A Facile and Straightforward Way to Synthesize Thiophene-Flanked Benzothiadiazole Derivatives and Their Polymers

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Figure S1. ¹H NMR (300 MHz) spectrum of DTBT in CDCl₃.



Figure S2. ¹³C NMR (300 MHz) spectrum of DTBT in CDCl₃.



Figure S3. ¹H NMR (300 MHz) spectrum of DHTBT in CDCl₃.







Figure S4. ¹³C NMR (300 MHz) spectrum of DHTBT in CDCl₃.



Figure S6. ¹H NMR (300 MHz) spectrum of PFTBT in C₂D₂Cl₄ at 100 °C.



Figure S7. ¹H NMR (300 MHz) spectrum of PFHTBT in C₂D₂Cl₄ at 100 °C.



Figure S8. ¹H NMR (300 MHz) spectrum of PFTBT-R in C₂D₂Cl₄ at 100 °C.



Figure S9. ¹H NMR (300 MHz) spectrum of PFHTBT-R in C₂D₂Cl₄ at 100 °C.



Figure S10. ¹H NMR (300 MHz) spectrum of PFEDOTBT-R in $C_2D_2Cl_4$ at 100 °C.



Figure S11. GPC curves of the alternating copolymers.



Figure S12. GPC curves of the random copolymers.



Figure S13. Comparison of normalized UV-Vis absorption spectra of the alternating polymer PFHTBT and the random polymer PFHTBT-R.