Electron-transporting Thiazole-based polymer synthesized through direct (hetero)arylation polymerization

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Supplementary Information.

1. Polymer ¹H NMR traces



Figure S1. ¹H NMR spectrum of the P(TzDPP-Th) batch B1 in CDCl₃



Figure S2. ¹H NMR spectrum of the P(TzDPP-Th) batch B2 in CDCl₃



Figure S3. ¹H NMR spectrum of the P(TzDPP-Th) batch B3 in C₂D₂Cl₄



Figure S4. ¹H NMR spectrum of the P(TzDPP-Th) batch B4 in C₂D₂Cl₄



2. Polymer SEC chromatograms

Figure S5. SEC chromatogram of the P(TzDPP-Th) batch B1



Figure S6. SEC chromatogram of the P(TzDPP-Th) batch B2



Figure S7. SEC chromatogram of the P(TzDPP-Th) batch B3



Figure S8. SEC chromatogram of the P(TzDPP-Th) batch B4

3. Cyclic voltammogram



Figure S9. Cyclic voltammogram of P(TzDPP-Th) reference in thin-film recorded in acetonitrile + 0.2 M [NBu₄][PF₆]. Cyclic voltammogram of ferrocene vs SCE. Platinum working electrode, sweep-rate: 100 mV.s⁻¹.

4. OFET output and transfer characteristic





Figure **S10**. Output characteristics of the OFETs whose channel is **B2** (a), **B3** (b) and **B4** (c). Ids is the drain-source current, Vds the voltage difference between drain and source and V_{gs} the voltage difference between gate and source.



Figure **S11**. Transfer characteristics measured on the same **B2**, **B3** and **B4** OFETs at a drain-source voltage difference (V_{ds}) of 80 V. The electron mobility is directly proportional to the slope of the square-root of I_{ds} as a function of V_{gs} using the standard formalism of OFETs in the saturation regime.