Supporting Information

Effects of pyridyl group orientations on the optoelectronic properties of regio-isomeric diketopyrrolopyrrole based π conjugated polymers

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	<i>M</i> n ^{<i>a</i>} (kDa)	Mw ^a (kDa)	PDI	Yield (%)	<i>T</i> _d ^b (°C)
d-PDBPy	37.7	94.1	2.5	82	413
p-PDBPy	21.4	67.5	3.1	69	394

Table S1. Molecular weight and thermal properties of the copolymers.

^{*a*} Determined by GPC in tetrahydrofuran (THF) using polystyrene standards.

^b The 5% weight-loss temperatures under a nitrogen atmosphere.



Fig. S1 (a) TGA curves of copolymers at a heating rate of 20 °C min⁻¹ under a nitrogen

atmosphere; (b) DSC characteristics of polymers at a scanning rate 10 °C min⁻¹.



Fig. S2 Output and transfer characteristics of (a and b) *d*-PDBPy and (c and d) *p*-PDBPy devices (spin-coated from CF solutions) at V_{SD} =-30 V (*L*=70 μ m, *W*=500 μ m) after thermal annealing at 100 for 10 min.



Fig. S3 The 500 MHz ¹H NMR spectrum of compound 1.



Fig. S4 The 126 MHz 13 C NMR spectrum of compound 1.



Fig. S5 The 500 MHz ¹H NMR spectrum of monomer 2.



Fig. S6 The 126 MHz ¹³C NMR spectrum of monomer 2.



Fig. S7 The 500 MHz ¹H NMR spectrum of monomer 4.



Fig. S8 The 126 MHz ¹³C NMR spectrum of monomer 4.



Fig. S9 The 500 MHz ¹H NMR spectrum of compound 5.



Fig. S10 The 126 MHz ¹³C NMR spectrum of compound 5.



Fig. S11 The 500 MHz ¹H NMR spectrum of compound 7.



Fig. S12 The 126 MHz ¹³C NMR spectrum of compound 7.



Fig. S13 The 500 MHz ¹H NMR spectrum of monomer 8.



Fig. S14 The 126 MHz ¹³C NMR spectrum of monomer 8.