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Supplementary Information

Organic Field Effect Transistors Based on Self-Assembling Core-Modified Peptidic Polymers

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Scheme S1: Synthetic route of diacetylene monomers F1 and F2.



Scheme S2: Synthesis of Phe-appended polydiacetylenes PF and PFF.

Polymer	HOMO vs. Vacuum	LUMO vs. Vacuum	Band Gap (From CV) (eV)	Band Gap (From UV) (eV)
PFF	-6.82	-4.50	2.32	2.24
PF	-6.75	-4.31	2.44	2.28

 Table S1: Details of the band structure parameters of polymers.



Figure S1: Concentration-dependent ¹H NMR (300 MHz, CDCl₃) of **F1**. (Concentration-dependent ¹H NMR spectra of **F2** was not recorded due to its immediate polymerization)



Figure S2: Normalized Raman spectra of diacetylenes and corresponding polydiacetylenes. Blue dotted line is diacetylene before UV exposure and black solid line is after the formation of polymers. (Raman spectrum of **F2** was not recorded due to its immediate polymerization)



Figure S3: UV-Vis absorption spectra of diacetylene monomers and corresponding polymers. (UV-Vis absorption spectrum of **F2** was not recorded due to its immediate polymerization)



Figure S4: Powder X-ray diffraction pattern of polymer PFF. (PXRD data of PF was not recorded due to its sticky nature)



Figure S5: FET measurements on polymer PF (a) Output characteristics, (b) Transfer characteristics.



Figure S6: Variation of mobility with V_{GS} at room temperature under ambient conditions in the case of PFF.



Figure S7: ¹H NMR (300 MHz, CDCl₃) spectrum of M1.



Figure S8: ¹³C NMR (75 MHz, CDCl₃) spectrum of M1.



Figure S9: ESI-Mass Spectrum of M1.



Figure S10: ¹H NMR (500 MHz, DMSO- d^6) spectrum of F1.



Figure S11: 13 C NMR (125 MHz, DMSO- d^6) spectrum of F1.



Figure S12: ESI-Mass Spectrum of F1.



Figure S13: ¹H NMR (300 MHz, CDCl₃) spectrum of M2.



Figure S14: ¹³C NMR (75 MHz, CDCl₃) spectrum of M2.



Figure S15: ESI-Mass Spectrum of M2.