Thermoresponsive dendronized poly(phenylacetylene)s showing tunable helicity

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Figure S1. ¹H NMR spectrum of monomer PA-ACe in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S2. ¹H NMR spectrum of monomer PA-ACm in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S3. ¹H NMR spectrum of monomer **PA-ES** in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S4. ¹H NMR spectrum of monomer PA-dAM in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S5. ¹H NMR spectrum of monomer PA-AM in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S6. ¹H NMR spectrum of compound PPA-ACe in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S7. ¹H NMR spectrum of compound **PPA-ACm** in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S8. ¹H NMR spectrum of compound PPA-ES in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S9. ¹H NMR spectrum of compound PPA-dAM in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S10. ¹H NMR spectrum of compound **PPA-AM** in d_6 -DMSO. The solvent peak is marked with asterisk (*).



Figure S11. Atomic force microscopy (AFM) image of **PPA-ACm** from THF solution (0.002 mg/mL).



Figure S12. Atomic force microscopy (AFM) image of **PPA-ACe** from THF solution (0.002 mg/mL).



Figure S13. CD and UV-vis spectra of **PPA-ES** (0.25 mg/mL) measured in H₂O, THF, MeOH, DCM and CHCl₃ at 25 °C.



Figure S14. ¹H NMR spectra of monomer PA-ACm at various temperatures in D_2O , as well as in d₆-DMSO at 25 °C.



Figure S15. ¹H-¹H NOESY spectrum of monomer PA-ACm in D_2O at 10 °C. Mixing time 1000 ms.



Figure S16. ¹H-¹H NOESY spectrum of monomer PA-ACm in D_2O at 38 °C. Mixing time 1000 ms.



Figure S17. Temperature variable CD spectra of PPA-dAM (0.025 mg/mL) taken after the addition of $BaCl_2 \cdot H_2O$ (0.25 mg/mL).