

Supporting information

Synthesis and Self-Assembly of Poly(ferrocenyldimethylsilane)- *block-poly(2-alkyl-2-oxazoline)* Block Copolymers

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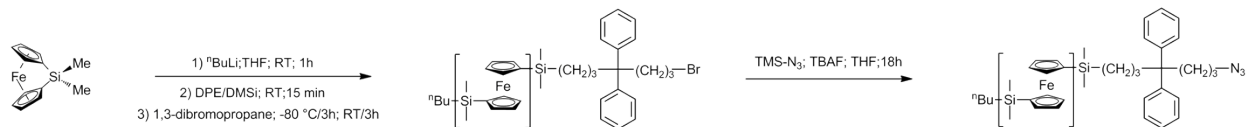


Figure S1: Synthesis of azide-functionalized PFDMS-N₃ of different molar mass.

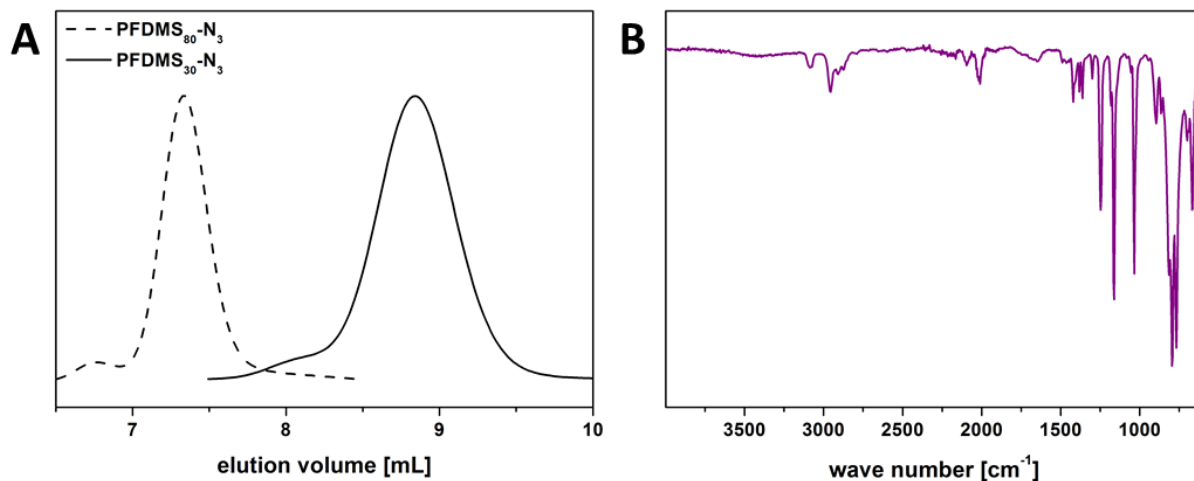


Figure S2: A) Comparison of the SEC traces for azide-modified PFDMS-N₃ with two different molar mass of 7 000 g mol⁻¹ (PFDMS₃₀-N₃; straight black line) and 20 000 g mol⁻¹ (PFDMS₈₀-N₃; dashed line); B) FT-IR spectrum for PFDMS₃₀-N₃.

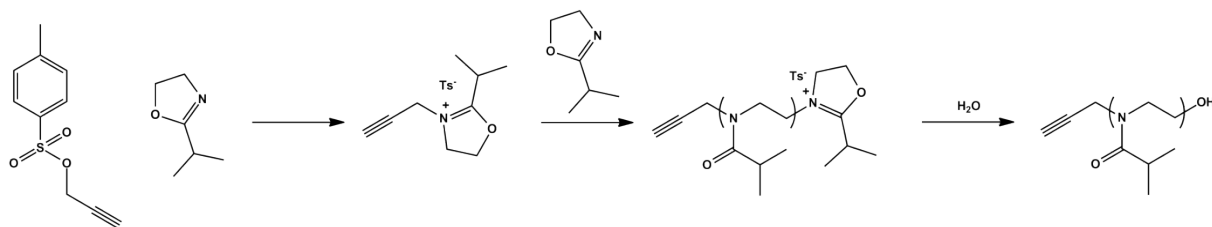


Figure S3: Synthesis of alkyne-functionalized PiPrOx *via* CROP initiated by propargyl *p*-toluenesulfonate.

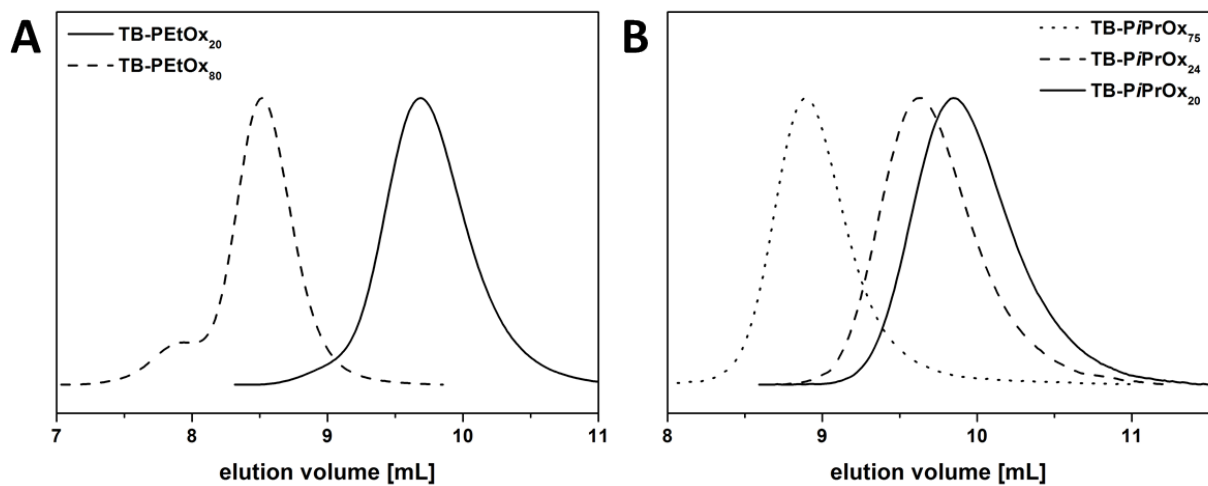


Figure S4: Comparison of SEC traces obtained for: A) TB-PEtOx₂₀ (straight line), TB-PEtOx₈₀ (dashed line); B) TB-PiPrOx₂₀ (straight line), TB-PiPrOx₂₄ (dotted line), and TB-PiPrOx₇₅ (dashed line).

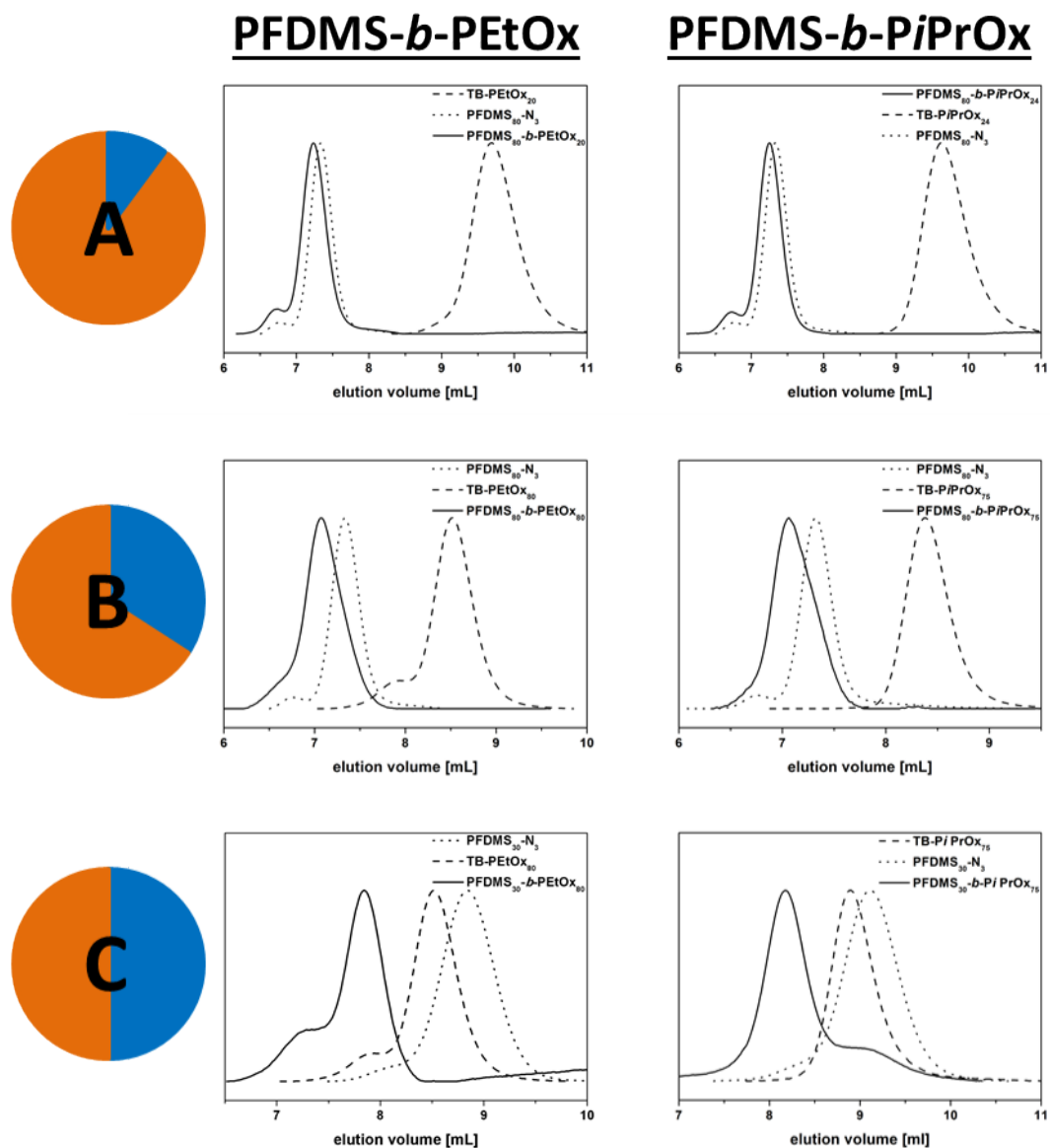


Figure S5: Comparison of SEC traces for the CuAAC click reaction between $\text{PFDMS}_x\text{-N}_3$ (dotted line) and various TB-PO_y (dashed line) synthesized block copolymers (straight line) with different ratios: A) $\text{PFDMS}_{80}\text{-}b\text{-PEtOx}_{20}$ (92/8 wt. %), $\text{PFDMS}_{80}\text{-}b\text{-PiPrOx}_{24}$ (88/12 wt. %); B) $\text{PFDMS}_{80}\text{-}b\text{-PEtOx}_{80}$ (71/29 wt. %), $\text{PFDMS}_{80}\text{-}b\text{-PiPrOx}_{75}$ (70/30 wt. %); C) $\text{PFDMS}_{30}\text{-}b\text{-PEtOx}_{80}$ (47/53 wt. %), $\text{PFDMS}_{30}\text{-}b\text{-PiPrOx}_{75}$ (46/54 wt. %).

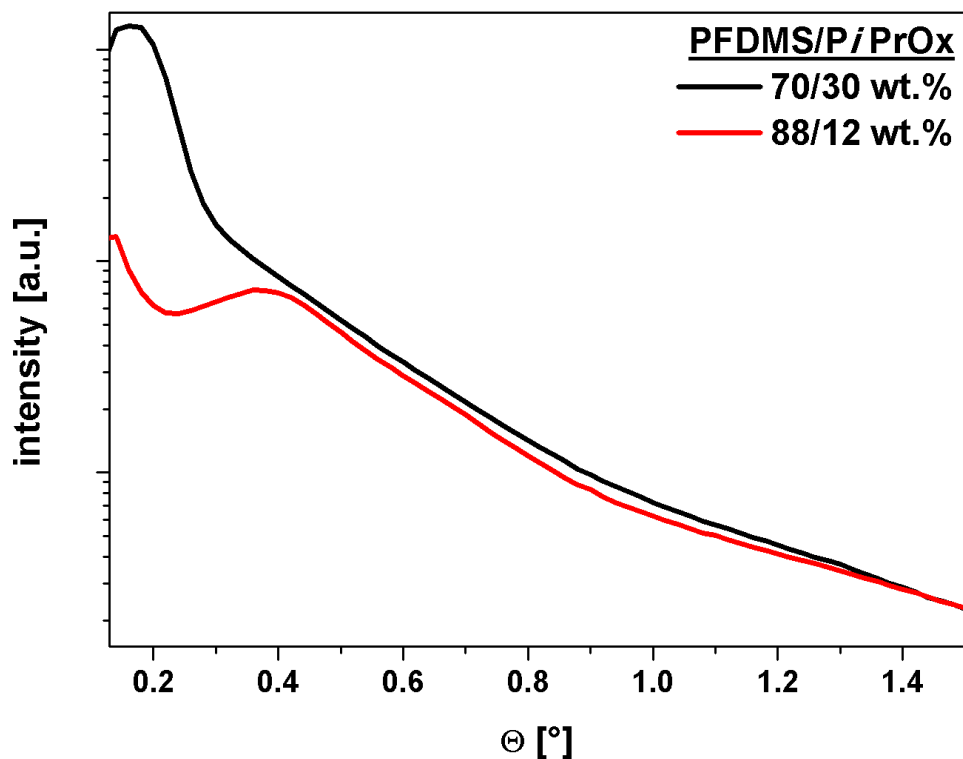


Figure S6: Comparison of SAXS patterns for different PFDMS_x-*b*-PiPrOx_y diblock copolymers: 70/30 wt. % (PFDMS₈₀-*b*-PiPrOx₇₅; black curve), 88/12 wt. % (PFDMS₈₀-*b*-PiPrOx₂₄; red curve).

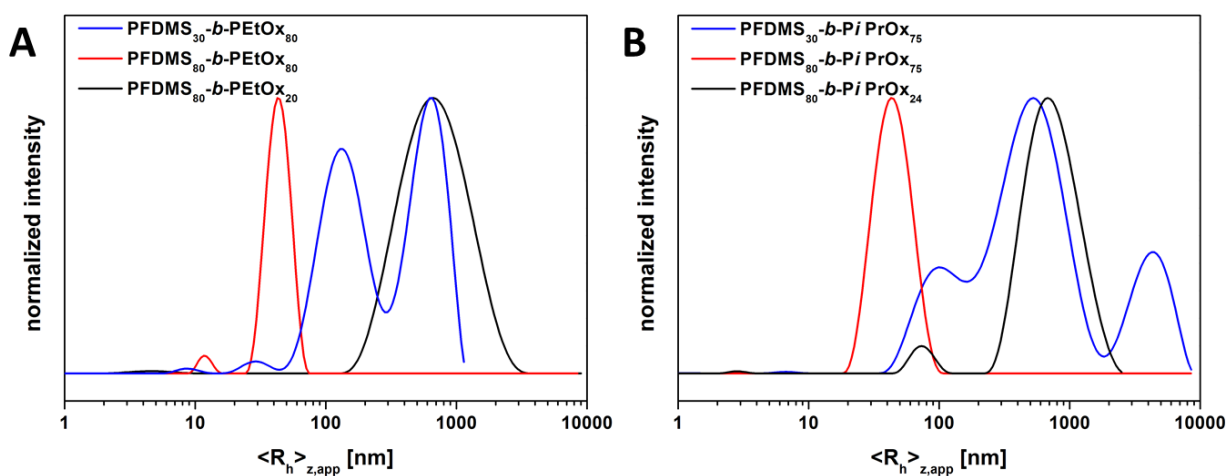


Figure S7: Comparison of DLS CONTIN plots for A) PFDMS₈₀-*b*-PEtOx₂₀ (92/8 wt. %; $\langle R_h \rangle_{z,app} = 670$ nm; black trace), PFDMS₈₀-*b*-PEtOx₈₀ (71/29 wt. %, $\langle R_h \rangle_{z,app} = 12$ and 43 nm; red trace), PFDMS₃₀-*b*-PEtOx₈₀ (47/53 wt. %; $\langle R_h \rangle_{z,app} = 9, 30, 135$ and 630 nm; blue trace); B) PFDMS₈₀-*b*-PiPrOx₂₄ (88/12 wt. %; $\langle R_h \rangle_{z,app} = 73$ and 680 nm; black trace); PFDMS₈₀-*b*-PiPrOx₇₅ (70/30 wt. %; $\langle R_h \rangle_{z,app} = 45$ nm; red trace); PFDMS₃₀-*b*-PiPrOx₇₅ (46/54 wt. %; $\langle R_h \rangle_{z,app} = 100, 525$ and 4300 nm; blue trace) at a constant concentration of 0.5 mg mL⁻¹ in acetone.

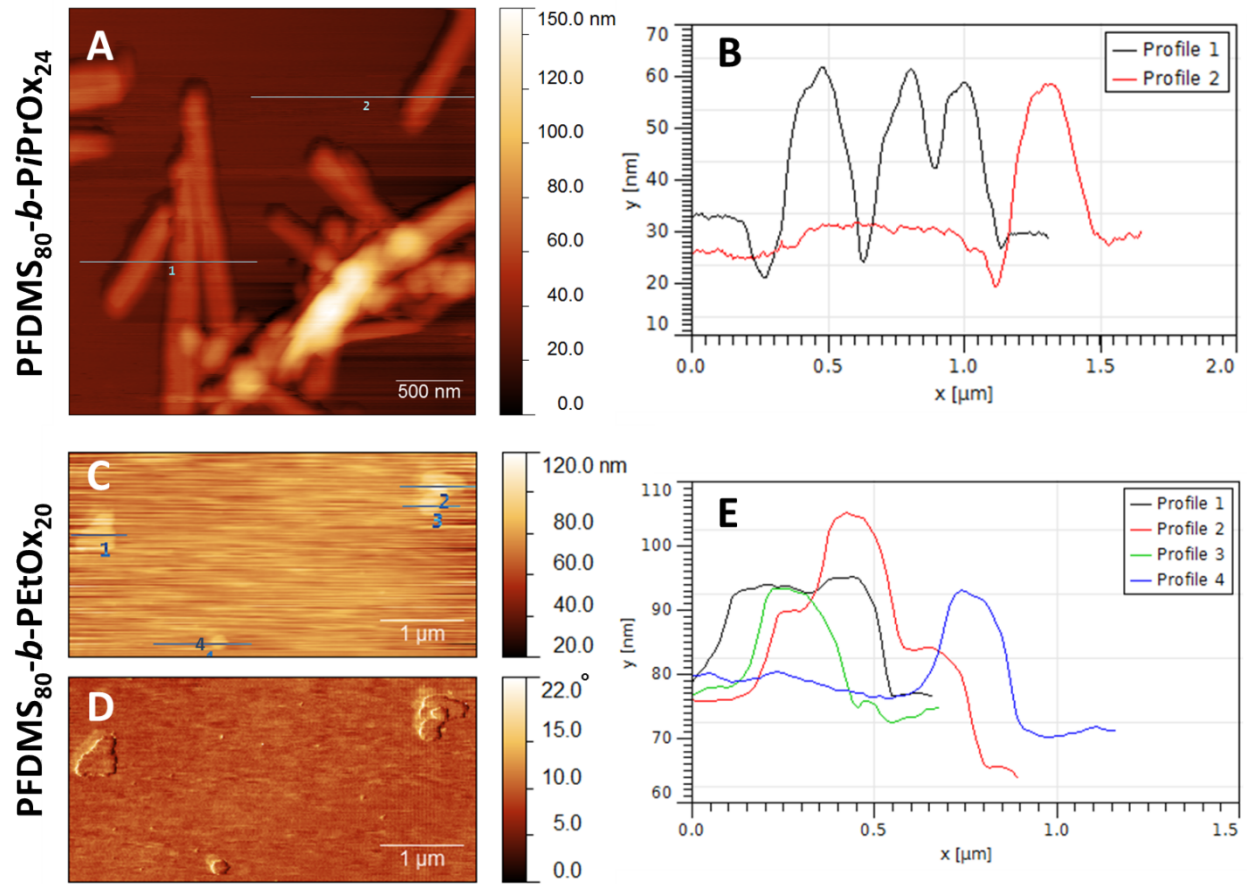


Figure S8: Comparison of AFM images for sheet forming block copolymers drop-casted from acetone: Height image (A) of PFDMS₈₀-*b*-PiPrOx₂₄ (88/12 wt. %) and the corresponding height profile (B, ~15-30 nm); height image (C) of PFDMS₈₀-*b*-PETox₂₀ (92/8 wt. %) and the corresponding phase image (D) and the height profile (E, ~15 nm).

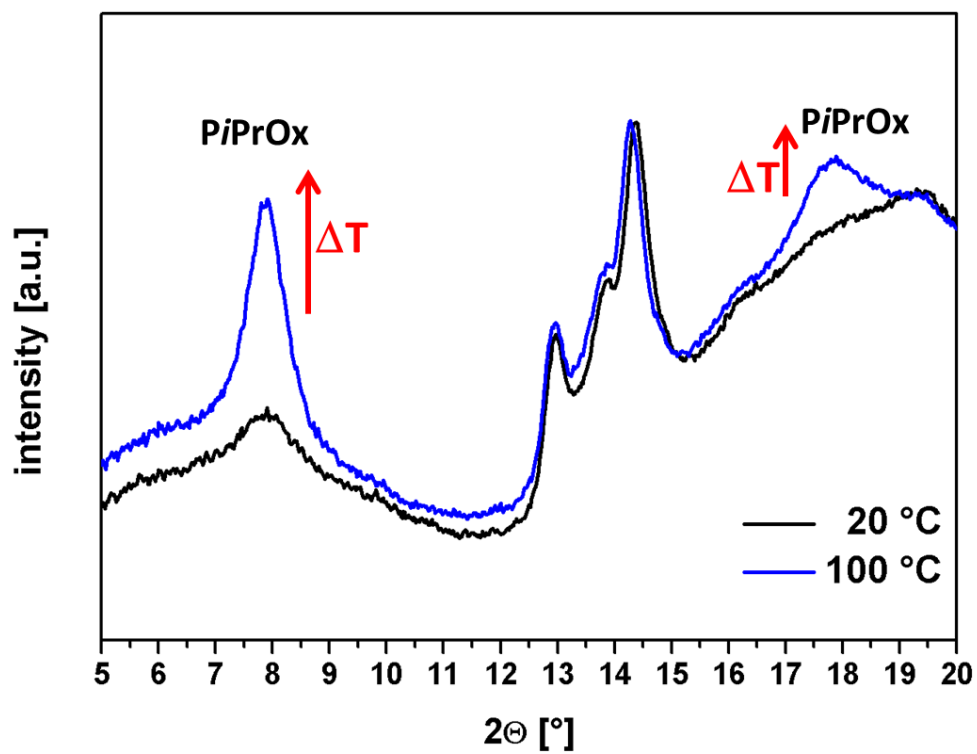


Figure S9: Wide angle x-ray scattering for PFDMS₃₀-*b*-PiPrOx₇₅ (46/54 wt.%) at different temperatures: 20 °C (black curve), and after heating to 100 °C (2 h@100 °C; blue curve).