Supporting Information for

Metallopolymer-Based Block Copolymers for the Preparation of Porous and Redox-Responsive Materials

Christian Rüttiger¹, Hanna Hübner¹, Sebastian Schöttner¹, Tamara Winter¹, Gennady Cherkashinin², Björn Kuttich³, Bernd Stühn³ and Markus Gallei^{1,*}

¹Ernst-Berl-Institute for Chemical Engineering and Macromolecular Science, Technische Universität Darmstadt, Alarich-Weiss-Str. 4, D-64287 Darmstadt, Germany

² Surface Science, Institute of Materials Science, Technische Universität Darmstadt, Jovanka-Bontschits-Str. 2, D-64287 Darmstadt, Germany

³Institute of Condensed Matter Physics, Technische Universität Darmstadt, Hochschulstraße 8, D-64289 Darmstadt, Germany

Corresponding author: m.gallei@mc.tu-darmstadt.de

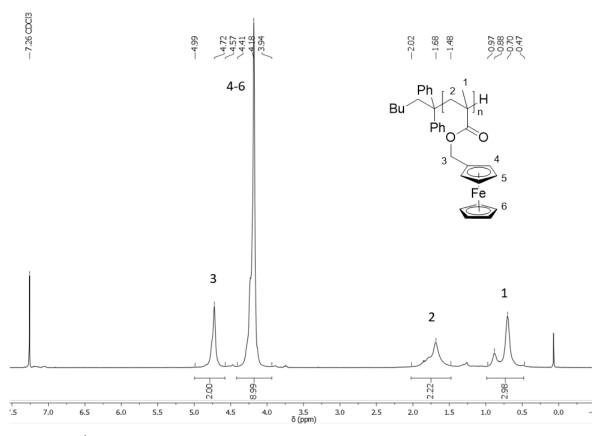


Figure S1. ¹H NMR spectrum of PFMMA₁₀₀ in CDCl₃.

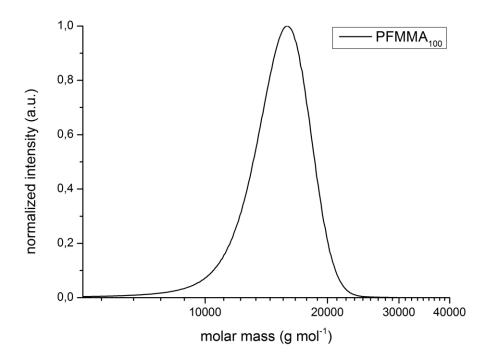


Figure S2. Molar mass distribution obtained by SEC measurements vs. PS standards in THF as eluent for PFMMA₁₀₀ (black line).

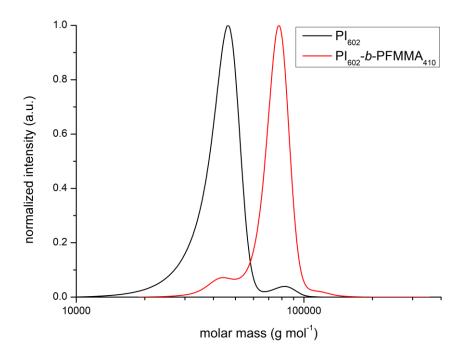


Figure S3. Molar mass distributions obtained by SEC measurements vs. PS standards in THF as eluent for PI₆₀₂ precursor (black line) and PI₆₀₂-*b*-PFMMA₄₁₀ (red line).

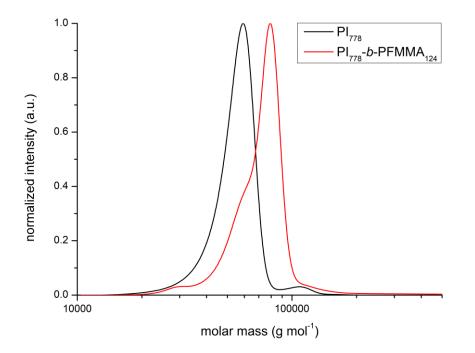


Figure S4. Molar mass distributions obtained by SEC measurements vs. PS standards in THF as eluent for PI₇₇₈ precursor (black line) and PI₇₇₈-*b*-PFMMA₁₂₄ (red line).

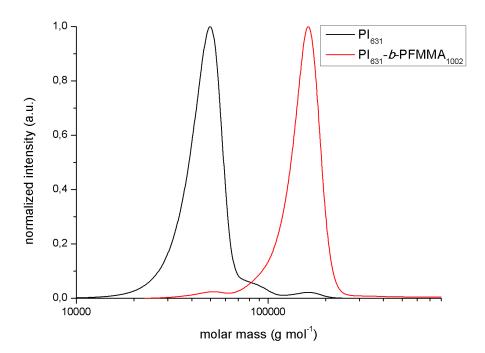


Figure S5. Molar mass distributions obtained by SEC measurements vs. PS standards in THF as eluent for PI₆₃₁ precursor (black line) and PI₆₃₁-*b*-PFMMA₁₀₀₂ (red line).

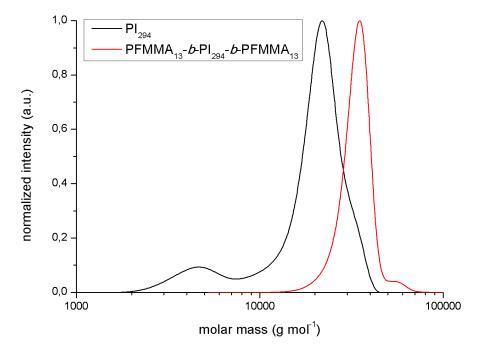


Figure S6. Molar mass distributions obtained by SEC measurements vs. PS standards in THF as eluent for PI₂₉₄ precursor (black line) and PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ (red line).

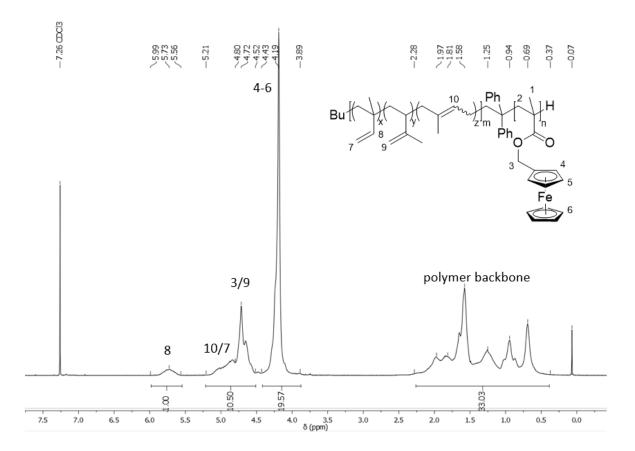


Figure S7. ¹H NMR spectrum of PI₆₀₂-*b*-PFMMA₄₁₀ in CDCl₃.

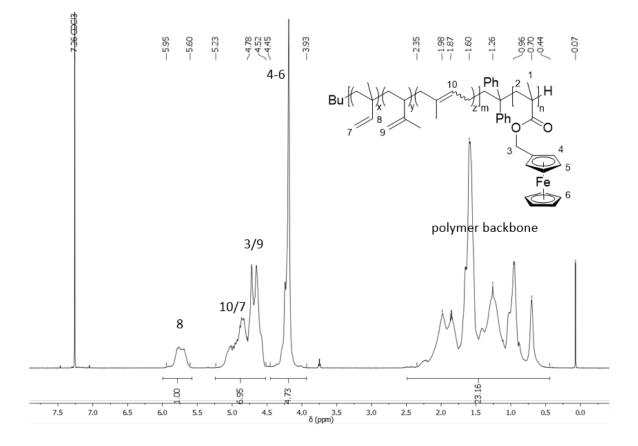


Figure S8. ¹H NMR spectrum of PI₇₇₈-*b*-PFMMA₁₂₄ in CDCl₃.

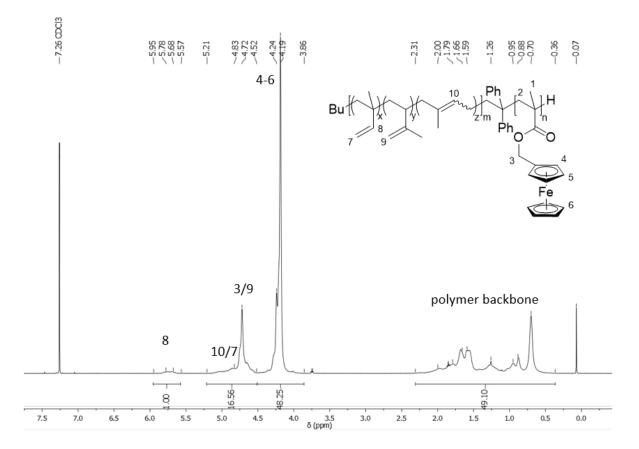


Figure S9. ¹H NMR spectrum of PI₆₃₁-*b*-PFMMA₁₀₀₂ in CDCl₃.

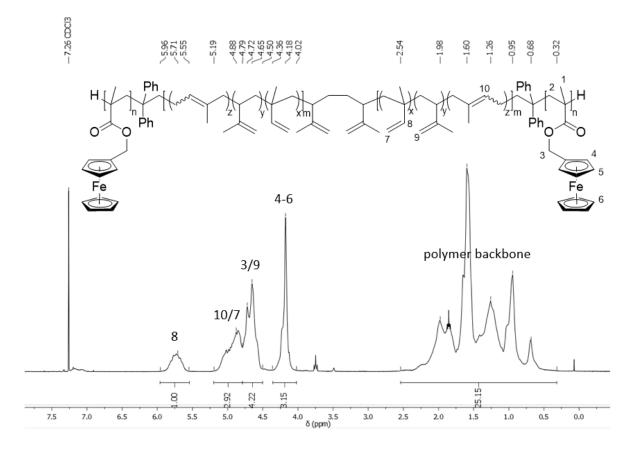


Figure S10. ¹H NMR spectrum of PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ in CDCl₃.

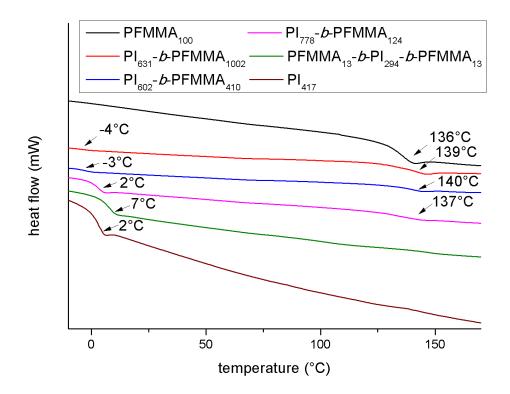


Figure S11. DSC thermograms of PFMMA₁₀₀ (black line), PI₆₃₁-*b*-PFMMA₁₀₀₂ (red line), PI₆₀₂-*b*-PFMMA₄₁₀ (blue line), PI₇₇₈-*b*-PFMMA₁₂₄ (pink line), PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ (green line) and PI₄₁₇ (brown line). The applied heat rate for the DSC run was 10 K min⁻¹ under nitrogen atmosphere.

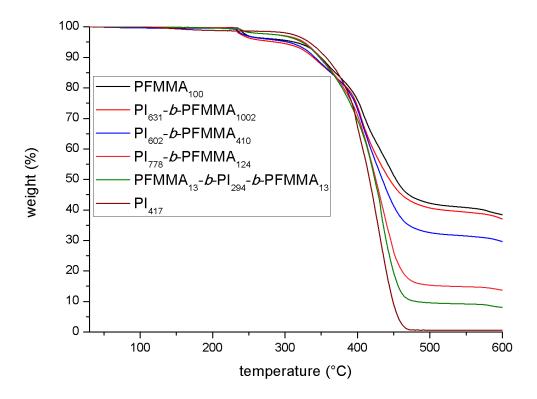


Figure S12. TGA curves of PFMMA₁₀₀ (black line), PI_{631} -*b*-PFMMA₁₀₀₂ (red line), PI_{602} -*b*-PFMMA₄₁₀ (blue line), PI_{778} -*b*-PFMMA₁₂₄ (pink line), PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ (green line) and PI_{417} (brown line). The applied heat rate for the TGA measurements was 10 K min⁻¹ in a nitrogen atmosphere until 600 °C.

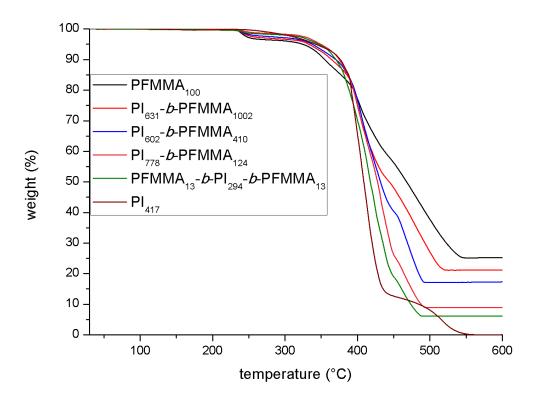


Figure S13. TGA thermograms of PFMMA₁₀₀ (black line), PI_{631} -*b*-PFMMA₁₀₀₂ (red line), PI_{602} -*b*-PFMMA₄₁₀ (blue line), PI_{778} -*b*-PFMMA₁₂₄ (pink line), PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ (green line) and PI_{417} (brown line). The applied heat rate for the TGA measurements was 10 K min⁻¹ in air atmosphere until 600 °C.

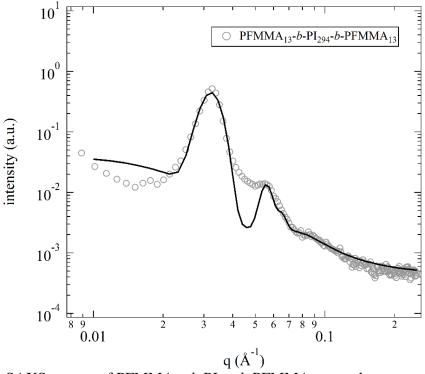


Figure S14. SAXS pattern of PFMMA₁₃-*b*-PI₂₉₄-*b*-PFMMA₁₃ sample.

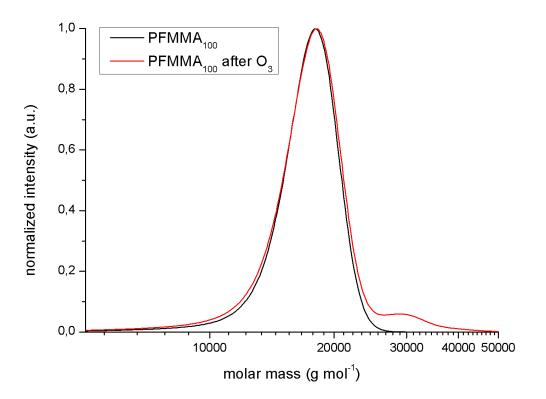


Figure S15. Molar mass distributions obtained by SEC measurements vs. PS standards in THF as eluent for PFMMA₁₀₀ (black line) and PFMMA₁₀₀ after two weeks in an ozone atmosphere (red line).

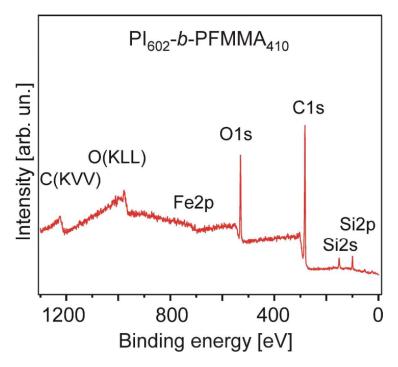


Figure S16. Al K α survey photoelectron spectrum of PI₆₀₂-*b*-PFMMA₄₁₀ deposited on a paper substrate prior to ozone treatment.

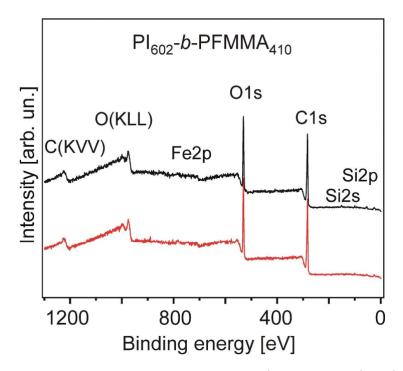


Figure S17. Al K α survey photoelectron spectra of PI₆₀₂-*b*-PFMMA₄₁₀ deposited on a paper substrate after ozone treatment measured at different areas of the sample.

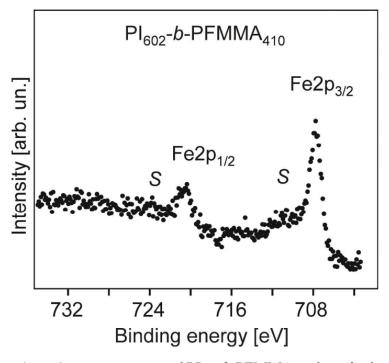


Figure S18. Fe2p photoelectron spectrum of PI₆₀₂-*b*-PFMMA₄₁₀ deposited on a paper substrate prior to ozone treatment.

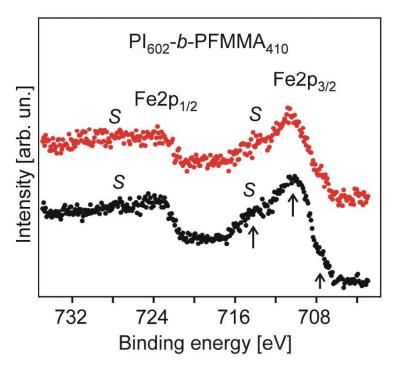


Figure S19. Fe2p photoelectron spectrum of sample PI_{602} -*b*-PFMMA₄₁₀ deposited on a paper substrate after ozone treatment measured at different areas of the sample.

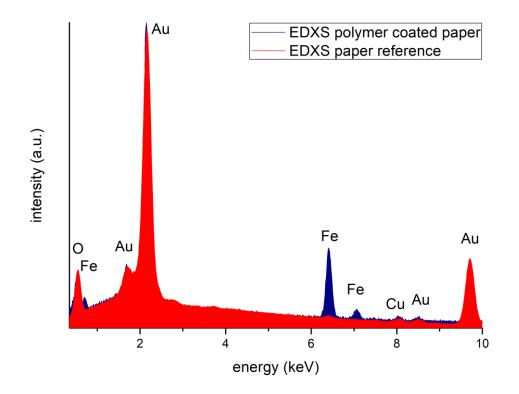


Figure S20. EDXS spectra of PI₆₀₂-*b*-PFMMA₄₁₀ coated paper substrate (blue) and untreated paper substrate (red) to prove the successful coating with synthesized ferrocene-containing polymer.

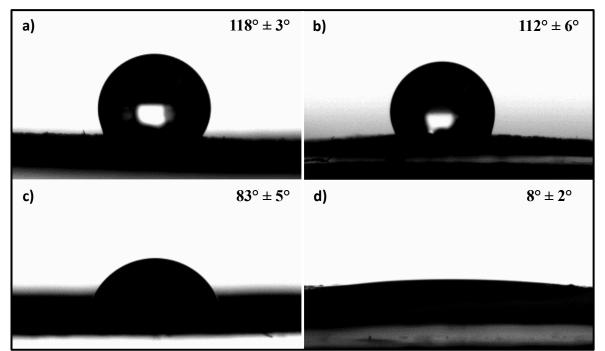


Figure S21. Contact angles obtained for PI_{602} -*b*-PFMMA₄₁₀ coated paper substrates: a) polymer coated paper substrate; b) polymer coated paper substrate after oxidation; c) polymer coated paper substrate after ozone and ethanol treatment; d) polymer coated paper substrate after ozone, ethanol treatment and oxidation.