



*Supplementary Information*

# Functionalizable Glyconanoparticles for a Versatile Redox Platform

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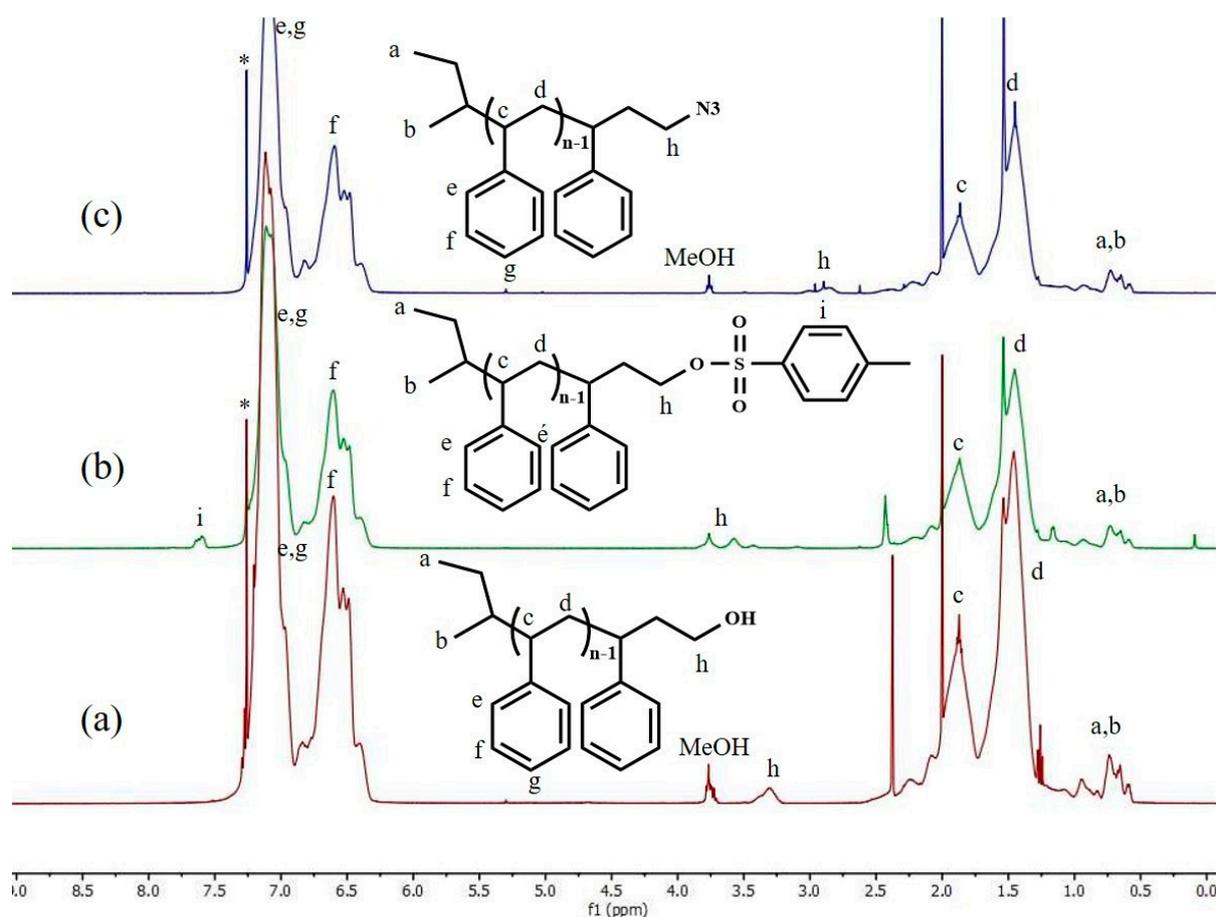


Figure S1.  $^1\text{H}$  NMR of (a) PS-OH, (b) PS-OTs and (c) PS-N<sub>3</sub> in  $\text{CDCl}_3$  at 25°C (400 MHz)

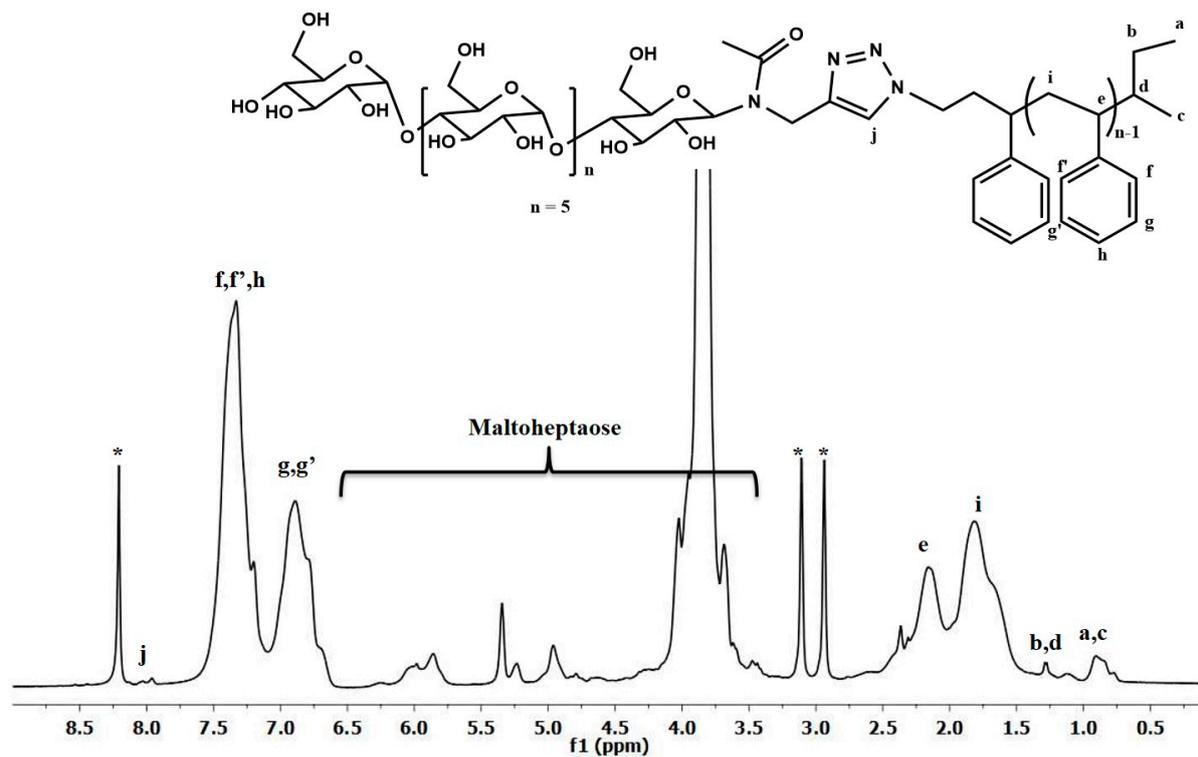
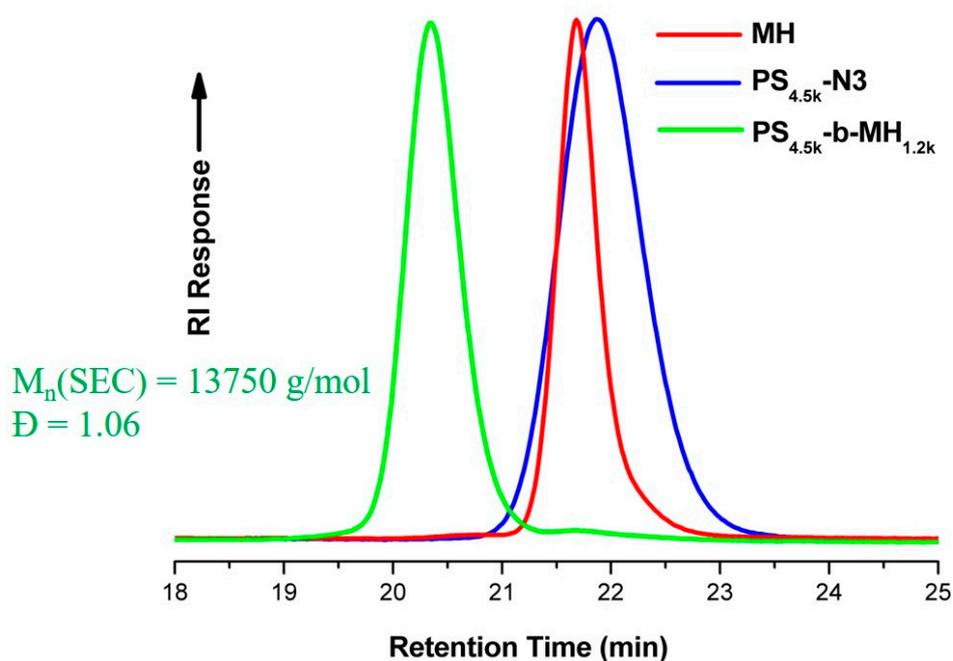
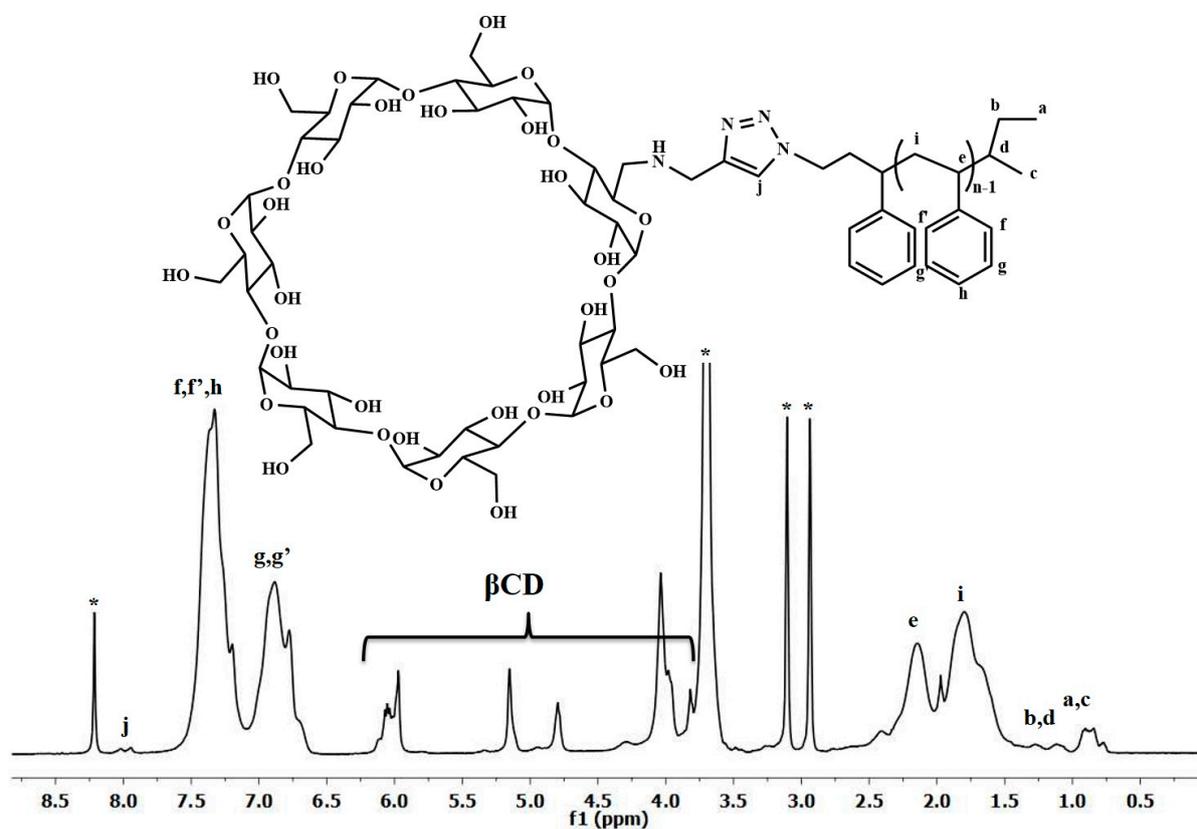


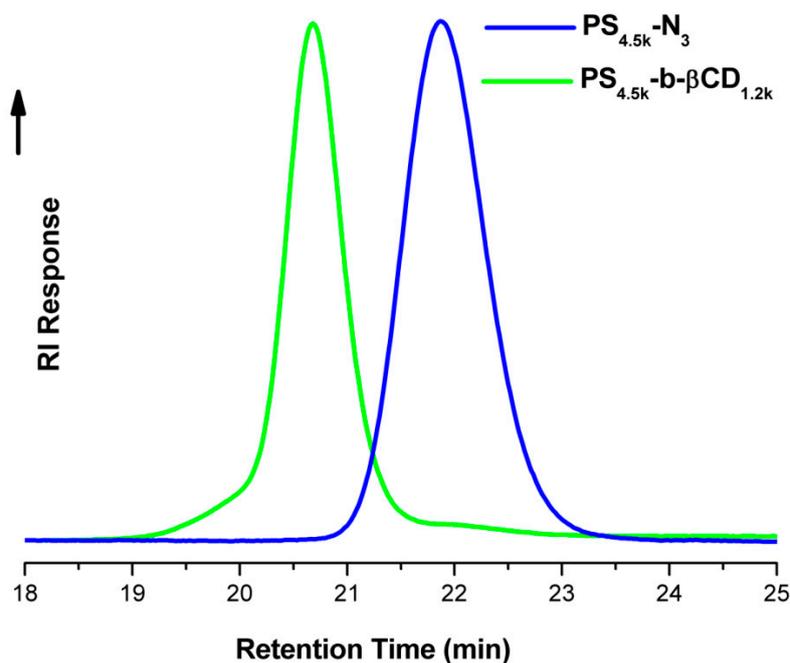
Figure S2.  $^1\text{H}$  NMR of PS-*b*-MH in  $\text{DMF-d}_7$  at 25°C (400 MHz)



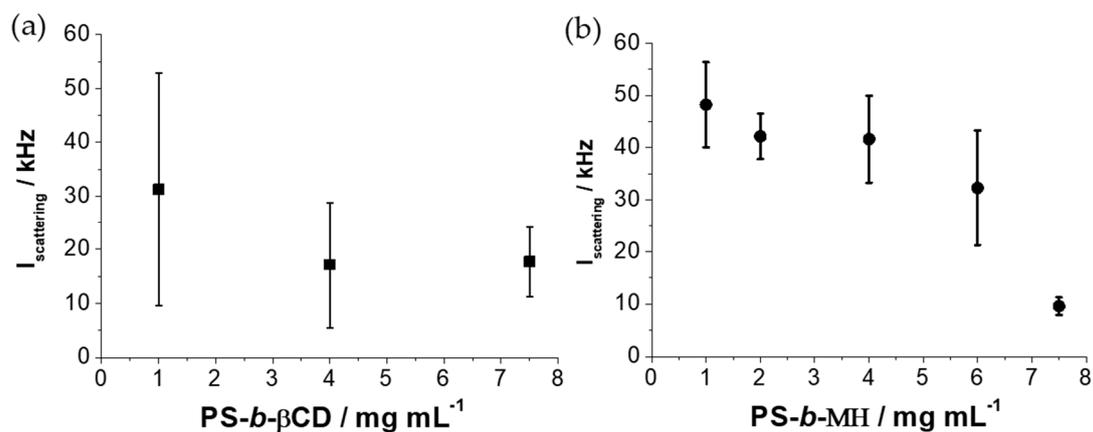
**Figure S3.** SEC traces of PS<sub>4.5k</sub>-N<sub>3</sub> (Blue), MH<sub>1.2k</sub> (Red) and PS<sub>4.5k</sub>-b-MH<sub>1.2k</sub> (green) using DMF as an eluent and PS calibration at 40°C.



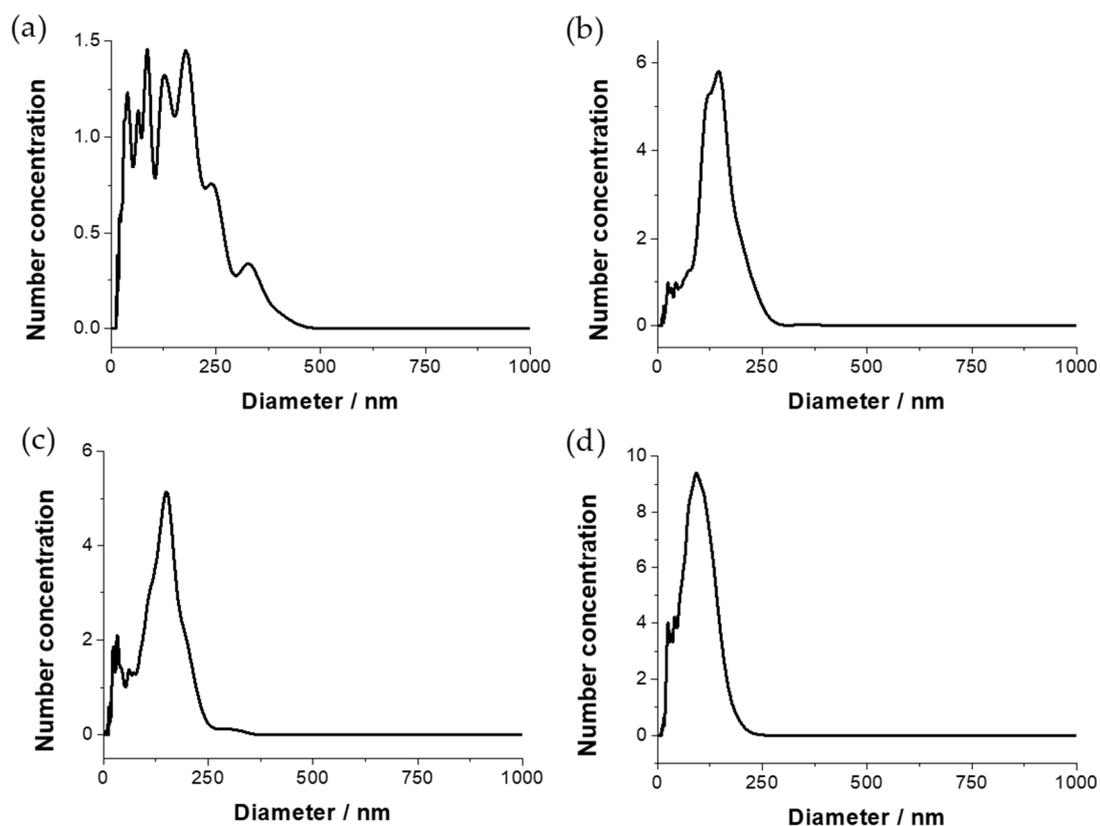
**Figure S4.**  $^1\text{H}$  NMR of PS-b- $\beta$ CD in DMF- $d_7$  at 25°C (400 MHz)



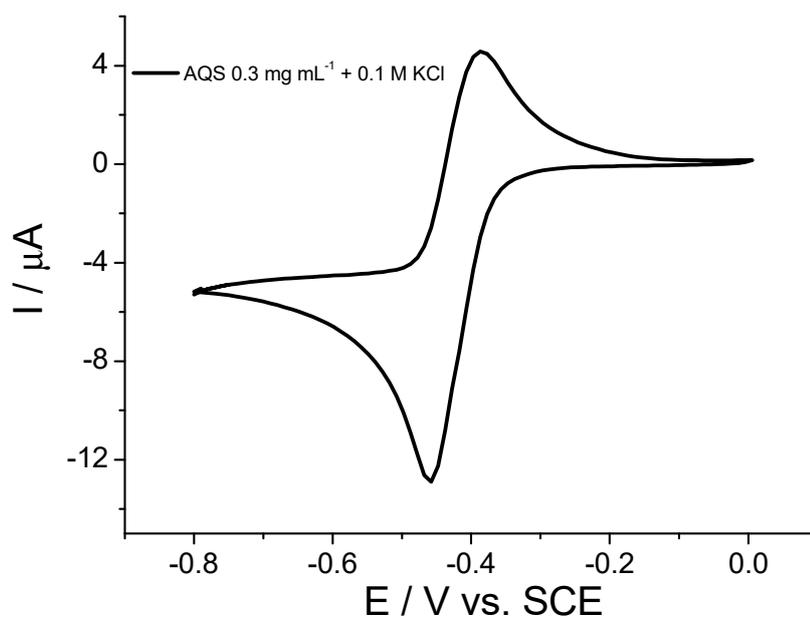
**Figure S5.** SEC traces of  $PS_{4.5k}-N_3$  (Blue), and  $PS_{4.5k}-b-\beta CD_{1.2k}$  (green) using DMF as an eluent and PS calibration at 40°C.



**Figure S6.** Scattering intensity as a function of mass concentration of  $PS-b-\beta CD$  and  $PS-b-MH$  glycopolymers in a THF/ $H_2O$  solution mixture (80:20 w/w %).



**Figure S7.** Size distribution of the solutions of (a) GNP<sub>PSCD</sub> (b) GNP<sub>PSMH</sub> (c) GNP<sub>PSCD50</sub> and (d) GNP<sub>PSCD10</sub> determined by NTA analysis.



**Figure S8.** Cyclic voltammetry performed at 10 mV s<sup>-1</sup> with glass carbon in aqueous solution (KCl 0.1 mol L<sup>-1</sup>) with AQS (0.3 mg mL<sup>-1</sup>). pH was adjusted to 6.0.