## Supporting Information

2-D Gold Nanoparticle Arrays from Thermally Directed Self-Assembly

of Peptide-Derivatized Block Copolymers

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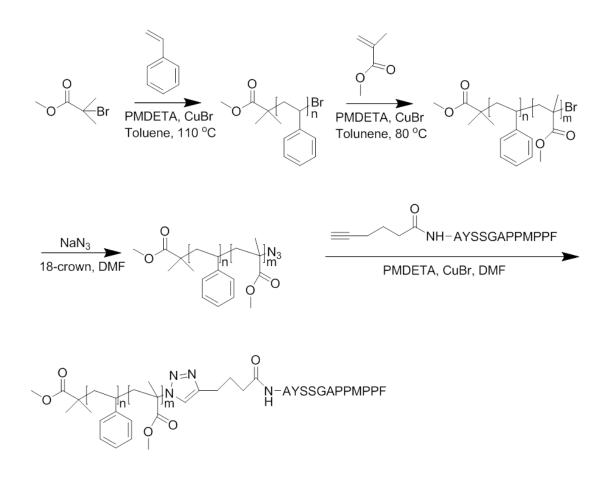


Figure S1. Synthetic route for A3 peptide-deviratized block copolymer bioconjugates. PS-b-PMMA was synthesized through a sequential polymerization followed by the "click" chemistry to tether A3 peptide at the end of PMMA.

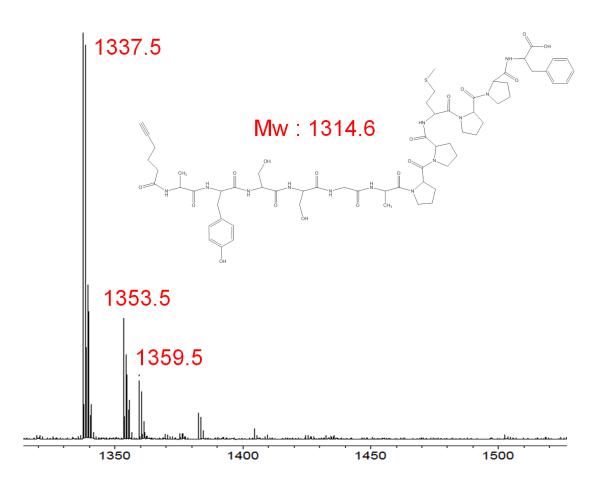


Figure S2. ESI-MS spectra of alkyne functionalized A3 peptide: 1337.5

(Alkyne-A3 peptide +Na<sup>+</sup>); 1353.5 (Alkyne-A3 peptide +Na<sup>+</sup>+K<sup>+</sup>-H<sup>+</sup>); 1359.5

(Alkyne-A3 peptide  $+2Na^{+}-H^{+}$ ).

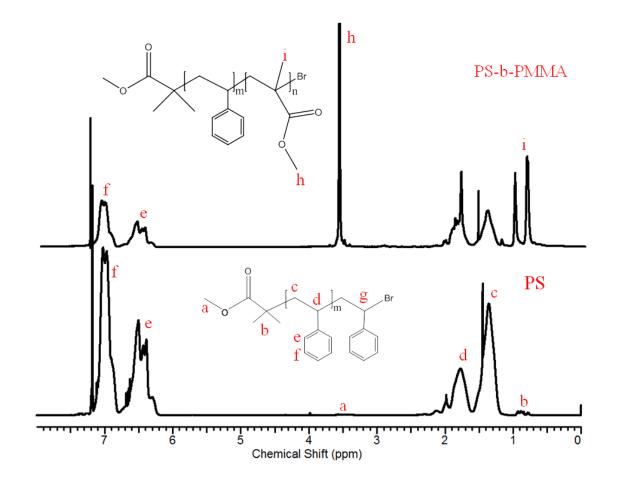


Figure S3. <sup>1</sup>H NMR (CDCl3, 500 MHz, ppm,  $\delta$ ) spectra of PS-Br (a) and PS-b-PMMA-Br (b). PS-Br was polymerized and used as the macromolecular initiator to grow PMMA block. PS-Br: 6.30-7.40 (br, 1980H, phenyl rings), 3.50 (s, 3H, CH<sub>3</sub>O-), 1.67-2.15 (br, 396H, -CH<sub>2</sub>CH(-Ar)-), 1.20-1.67 (br, 792H, -CH<sub>2</sub>CH(-Ar)-), 0.93 (s, 6H, -(C=O)C(CH<sub>3</sub>)<sub>2</sub>-). PS-b-PMMA-Br: 6.30-7.40 (br, 1980H, phenyl rings), 3.35-3.60 (br, 522H, CH<sub>3</sub>O-), 1.67-2.15 (br, 396H, -CH<sub>2</sub>CH(-Ar)-), 1.20-1.67 (br, 792H, -CH<sub>2</sub>CH(-Ar)-), 1.20-1.67 (br, 792H, -CH<sub>2</sub>CH(-Ar)-), 1.20-1.67 (br, 522H, CH<sub>3</sub>O-), 1.67-2.15 (br, 396H, -CH<sub>2</sub>CH(-Ar)-), 1.20-1.67 (br, 522H, -CH<sub>2</sub>CH(-Ar)-), 0.60-1.00 (br, 522H, -(CH<sub>3</sub>-)C(C=O) -).

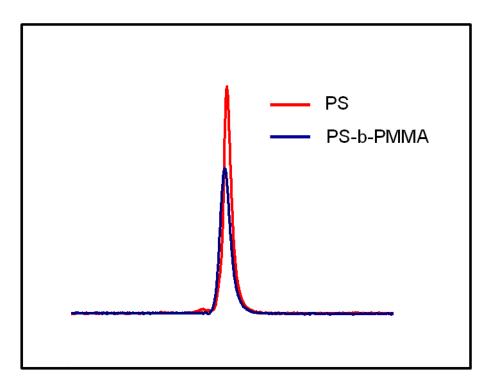


Figure S4. SEC spectra (THF, RI detector, PS standard) of PS-Br and PS-b-PMMA-Br. PS-Br was polymerized and used as the macromolecular initiator to grow PMMA block. PS-Br: Mn = 42.1 kDa, Mw = 43.9 kDa, PDI = 1.02. PS-b-PMMA-Br: Mn = 59.5 kDa, Mw = 69.0 kDa, PDI = 1.16.

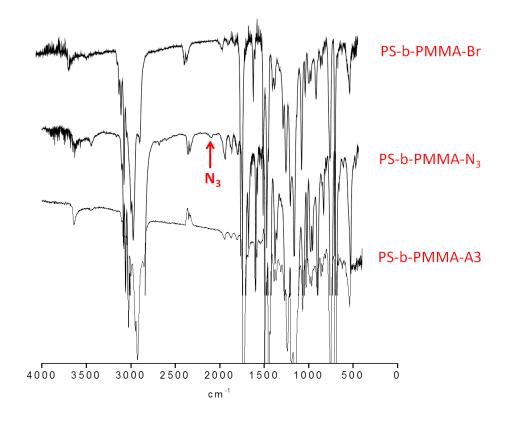


Figure S5. Fourier transform infrared (FTIR) spectra of PS-b-PMMA-Br, PS-b-PMMA-N<sub>3</sub> and PS-b-PMMA-A3. N<sub>3</sub> functional group showed signal at 2100  $cm^{-1}$  and disappeared after "Click" reaction.