A Highly Antibacterial Polymeric Hybrid Micelle with Efficiently Targeted Anticancer siRNA

Delivery and Anti-infection in vitro/in vivo

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Fig. S1. Synthesis routine of prepolymers and EHE copolymers. The reaction of P3/4HB with 1,4butylene glycol under catalytic PTSA to produce hydroxyl-terminated prepolymers (P3/4HB-diol) (A); PEG-OH coupling with HDI to form PEG-isocyanated (B); PEG-isocyanated coupling with P3/4HB-diol to form PEG-*b*-P3/4HB-OH (C); Prepolymer PEG-*b*-P3/4HB-OH coupling with acryloyl chloride to form PEG-*b*-P3/4HB-acrylated (D); PEG-*b*-P3/4HB-*b*-EPL (EHE) copolymer was synthesized by a Michael addition (E).









Fig. S2. NMR of P3/4HB-diol (A), PEG-P3/4HB-OH (B), PEG-P3/4HB-acrylated (C) in CDCl₃ EHP in H₂O (D) and EHP-FA in DMSO (E).



Fig. S3. Particle size and zeta potential of polymers without siRNA.



Fig. S4. Size distribution by intensity and raw correlation data of polymers without siRNA.



Fig. S5. Size distribution by intensity and raw correlation data of polymer/siRNA complexes.



Fig. S6. (A-C) Flow cytometric histogram profiles of Cy3 fluorescence intensity for cells with or without pre-treated by FA transfected by different polymer/Cy3-siRNA complexes. (D). Confocal microscopy image of cells treated with PEI/Cy3-siRNA complex.