Supporting Information

Size-Tailored and Acid-Degradable Polyvinyl Alcohol Microgels for Inhalation Therapy of Bacterial Pneumonia

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Figure S1. Preparation and characterization and of VEMA, PVA_{VEMA} and Cy5-PVA_{VEMA}. (A) Synthesis steps of VEMA. (B) Synthesis steps of PVA_{VEMA} and Cy5-PVA_{VEMA}. (C) ¹H NMR spectra of VEMA (400 MHz, CDCl₃). (D) ¹H NMR spectra of PVA_{VEMA} (400 MHz, DMSO-*d6*).



Figure S2. Size distribution of PVA_{VEMA} dispersed in water at various concentrations and at 5 °C (A), 10 °C (B) and 15 °C (C).



Figure S3. Particle size distribution of 0.5 mg/mL aqueous solution of PVA_{VEMA} aggregation and PVA_{VEMA} microgels (G) at 5 and 40°C. (Cross-linked: red dashed line, uncross-linked: black dashed line)



Figure S4. Size distribution of the PVA $_{VEMA}$ microgels before and after loading with PIP. (Before loading: red dashed line, After loading: solid line)



Figure S5. The pH degradation of PVA_{VEMA} microgel. (A) Gas chromatographic detection of PVA_{VEMA} microgel after complete degradation under acidic conditions. (B) Gas chromatographic detection of degradation of PVA_{VEMA} microgel at pH 7.4 and 6.6 for 120 h.



Figure S6. Antibacterial activity of the antibiotic drug PIP *in vitro*. (A) Coated plate method to evaluate the effect of different concentrations of PIP on *E. coli-lux* inhibition (A) and corresponding statistics (B). All data are presented as the mean \pm SD (n = 3).



Figure S7. *In vitro* analysis of antibacterial properties. (A) Changes in colony growth of *E. coli-lux* with different groups (Control, G-0.2@PIP, G-1.6@PIP, G-3.5@PIP and G-7.8@PIP, PIP: 1.8 mg mL⁻¹, MBC) incubated under pH 7.4 and pH 6.6 conditions. Survival statistics of the corresponding bacteria under conditions (B). All data are presented as the mean \pm SD (n = 3).



Figure S8. Modeling of pneumonia. (A) *In vitro* photograph of isolated lung organs 48 h after injection of different volumes of *E. coli-lux* solution into the right lung parenchyma. (B) Photographs of fluorescence imaging of living mice at the indicated time points after injection of *E. coli-lux* solutions (10 μ L/mouse) into the right lung parenchyma. All data are presented as the mean ± SD (n = 3).