

SUPPLEMENTARY INFORMATION

Selective isolation of *E. coli* associated with urinary tract infection using anti-fimbrial modified magnetic reduced graphene oxide nanoheaters

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Fabrication of chitosan modified magnetic particles (MP_{CS})

FeCl₂×4H₂O (0.34 g, 1.7 mmol) and FeCl₃×6H₂O (0.95 g, 3.5 mmol) were dissolved in deaerated water (20 mL) and subsequently added to a nitrogen-protected three-necked flask under sonication. The resulting mixture was heated at 50°C for 30 min. Then concentrated ammonium hydroxide (2 mL) was added dropwise and kept at 50°C for 30 min. The system was finally cooled to room temperature and the solid product was isolated *via* a non-uniform magnetic field generated by a Nd–Fe–B permanent magnet. The resulting Fe₃O₄ particles (MP) were washed six times with Milli-Q water to remove unreacted chemicals and then stored in water.

A water dispersion of bare MP (10 mg mL⁻¹, 1 mL) was mixed with chitosan (20 mg) and sonicated for 1 h at room temperature. The formed magnetic particles (MP_{CS}) were isolated by means of magnet and purified through six consecutive wash/precipitation cycles with water to ensure complete removal of unreacted chitosan. The precipitate was dried in an oven at 50°C.

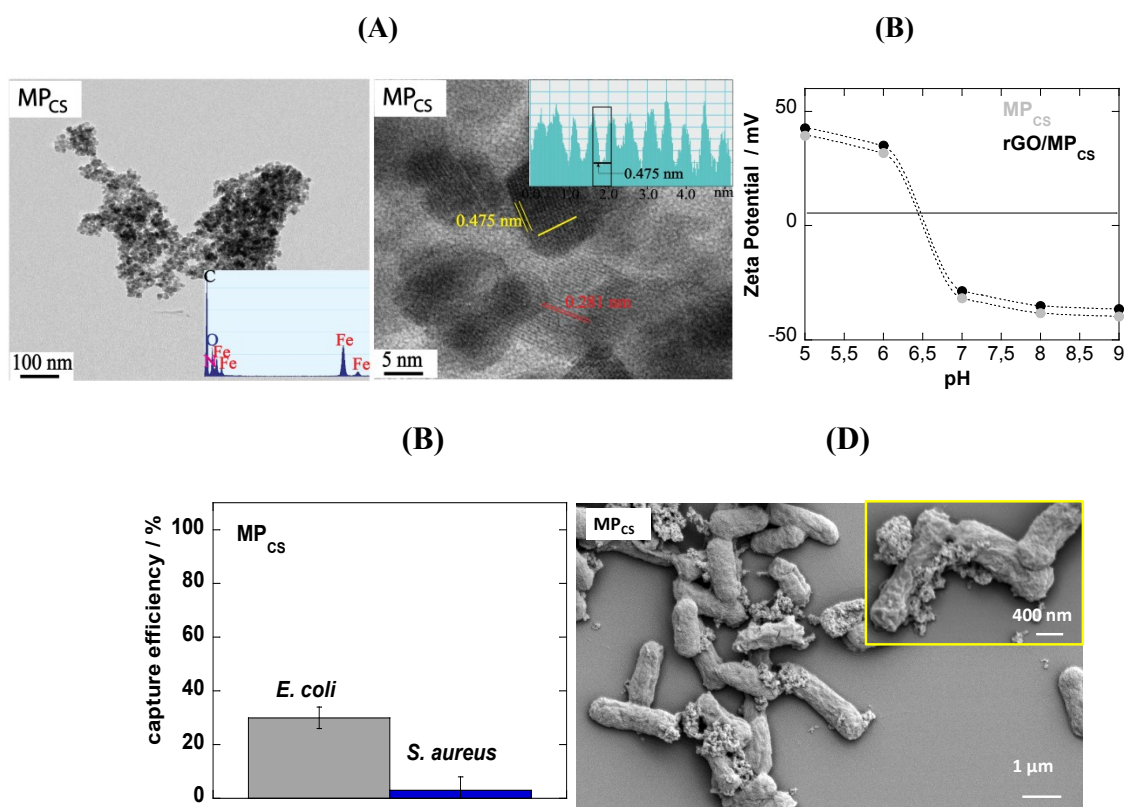


Figure S1: (A) Transmission electron microscopy images and HRTEM analysis of chitosan modified particles (Inset: EDAX analysis displaying the different elements present in the sample), (B) Zeta potential of MP_{CS} and rGO/MP_{CS} as a function of solution pH; (C) Removal efficiency of rGO/MP_{CS} (500 μg mL⁻¹) for *E. coli* and *S. aureus* (1×10⁹ cfu mL⁻¹); (D) SEM images of MP_{CS} nanoparticles mediated bacteria isolation.