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Electronic Supplementary Information (ESI)

Controllable Construction of Micro/nanostructured NiO in Confined

Microchannels via Microfluidic Chemical Fabrication for Highly Efficient and

Specific Absorption of Abundant Proteins

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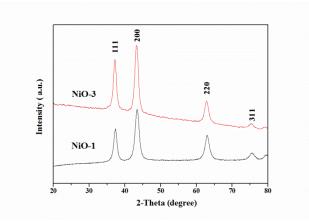


Fig. S1 XRD patterns of the as-prepared NiO scraped off the inner surface of capillary microchannels after calcined at 500 °C for 2 h.

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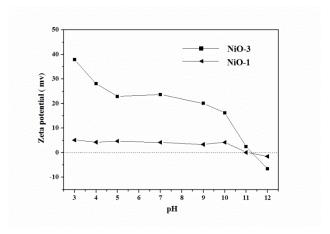


Fig. S2 Zeta-potential plots of the as-prepared NiO after calcined at 500 °C for 2 h. as a function of the pH of buffer solution.

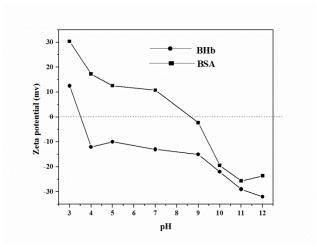


Fig. S3 Zeta-potential plots of BSA and BHb as a function of the pH of the buffer solution.

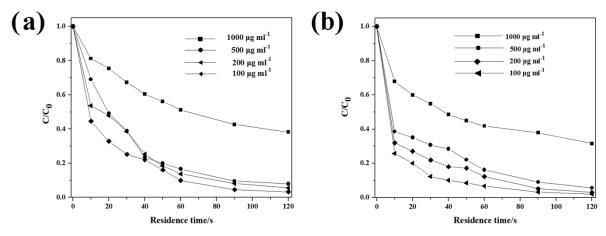


Fig. S4 Adsorption capture of proteins with different concentrations by NiO-3-modified capillary microchannels versus different RTs; (a) BSA; (b) BHb.

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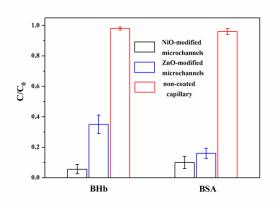


Fig. S5 Absorption capture of proteins (500 μg mL⁻¹) with NiO-3-modified microchannels, ZnO-modified microchannels and non-coated capillary.

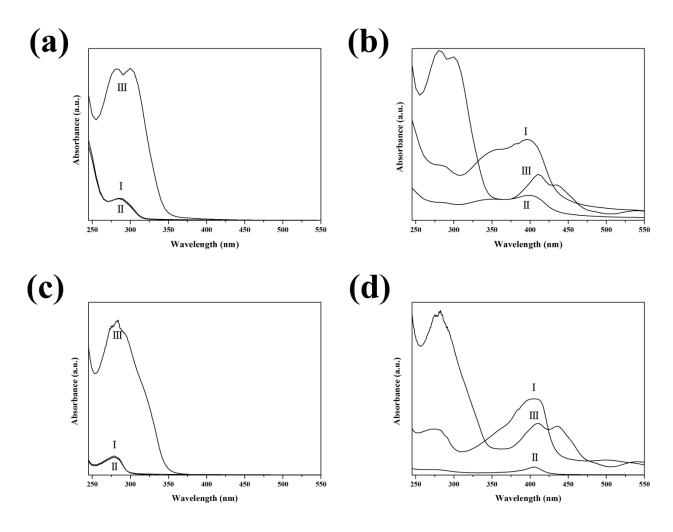


Fig. S6 UV-Vis spectra of the samples obtained in absorption-desorption process of single protein solution (500 μ g mL⁻¹) under two different conditions. (a) BSA, 1 mM KCl, pH=12; (b) BHb, 1 mM KCl,

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pH=12; (c) BSA, 200 mM KCl, pH=7; (d) BHb, 200 mM KCl, pH=7. Curve "I" "II" in the each map is the UV-Vis spectra of the samples before absorption, after absorption and after elution with $0.1~g~mL^{-1}$ imidazole solution respectively.