Supporting Information for

Polymer-coated NaYF₄:Yb³⁺, Er³⁺ Upconversion Nanoparticles for Charge-dependent Cellular Imaging

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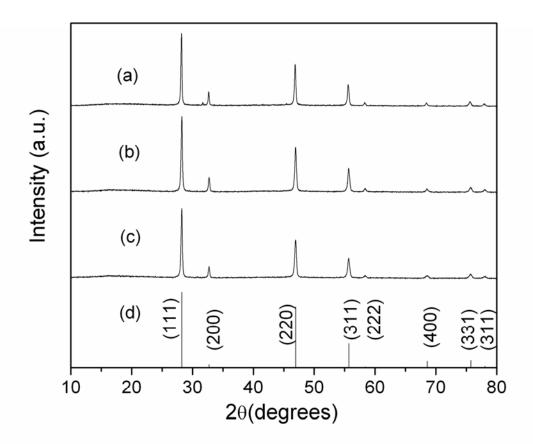


Figure S1. X-ray power diffraction (XRD) spectra of $NaYF_4:Yb^{3+}$, Er^{3+} NPs with various polymer coatings (a) UCNP-PEI, (b) UCNP-PVP, (c) UCNP-PAA, (d) Line pattern of the calculated cubic phase of $NaYF_4$ (PDF 77-2042).

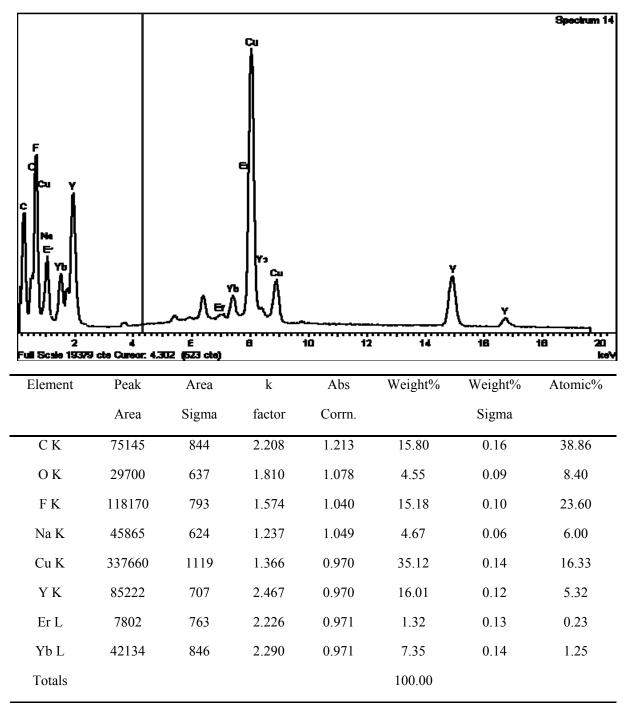


Figure S2. EDX spectrum of UCNP-PVP (upper) and the corresponding quantitative analysis as shown in the table list (lower).

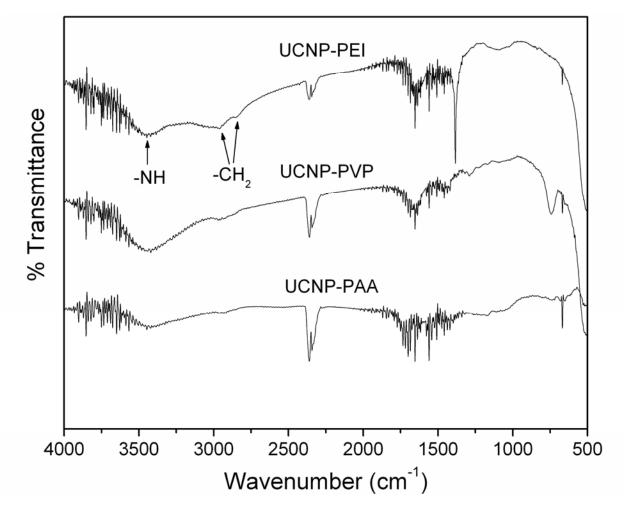


Figure S3. FT-IR spectra of NaYF₄:Yb³⁺, Er³⁺ NPs with various polymer coatings.

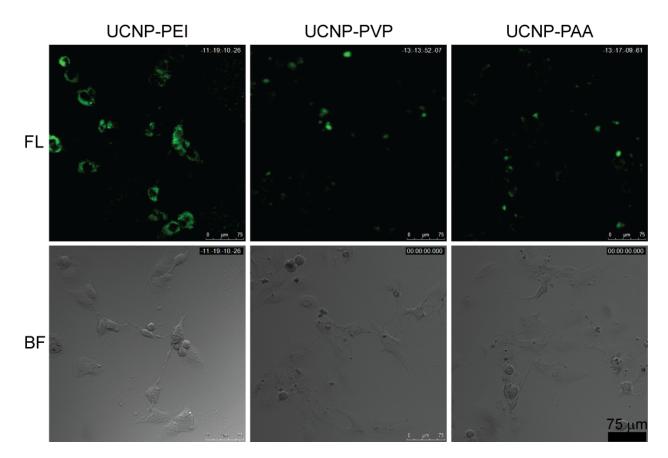


Figure S4. Multiphoton confocal fluorescent (upper) and bright field (lower) images of U87MG cells incubated with 50 μ g/ml UCNP-PEI (left panel), UCNP-PVP (middle panel) and UCNP-PAA (right panel). The 980 nm excitation was provided by a femto-second Ti:sapphire pulsed laser, and the green emissions of 540–560 nm were acquired by a PMT channel (10× lens, scale bar = 75 μ m).

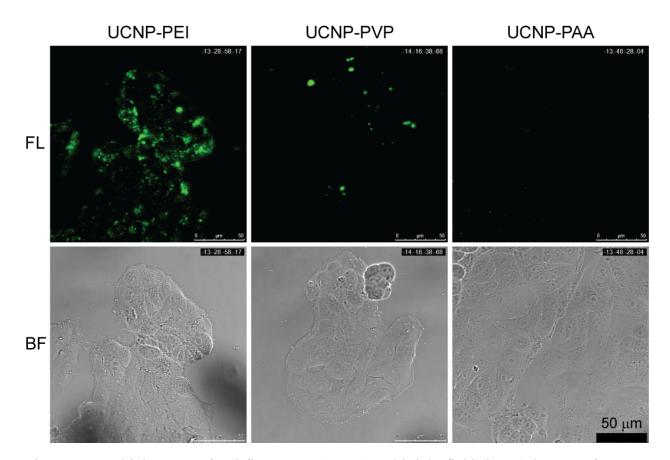


Figure S5. Multiphoton confocal fluorescent (upper) and bright field (lower) images of MCF-7 cells incubated with 50 μ g/ml UCNP-PEI (left panel), UCNP-PVP (middle panel) and UCNP-PAA (right panel). The 980 nm excitation was provided from a femto-second Ti:sapphire pulsed laser, and the green emissions of 540–560 nm were acquired by a PMT channel (40× oil lens, scale bar = 50 μ m).

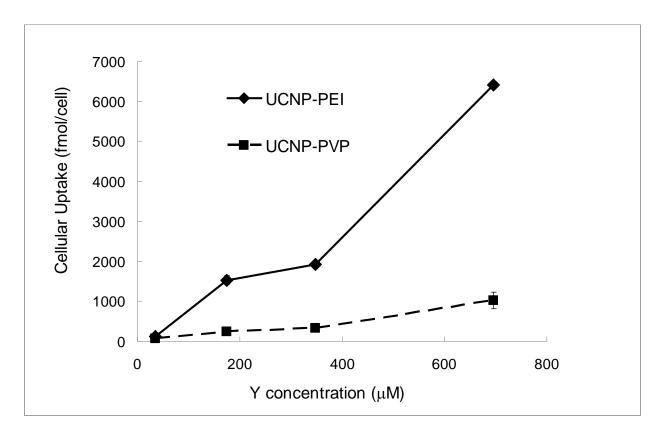


Figure S6. Average molar numbers of yttrium taken by an U87MG cell (fmol/cell) were determined by ICP-MS. U87MG cells were treated with the polymer coated UCNPs with yttrium concentrations ranging from 50 μ M to 400 μ M at 37 °C for 24 h. Each data point was represented as mean ± SD from triplicate trials.

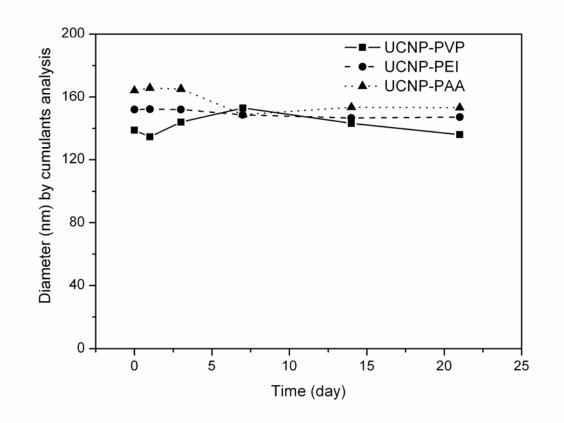


Figure S7. Long term stability test of UCNP-PVP (solid line), UCNP-PEI (dashed line) and UCNP-PAA (dotted line) with a dispersion concentration of 1 mg/mL in aqueous solution.

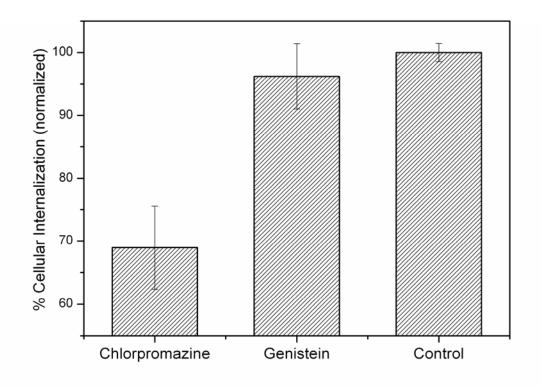


Figure S8. Probing the uptake pathways of UCNP-PEI using chemical endocytic inhibitors. HeLa cells were incubated with chlorpromazine and genistein for the inhibition of clathrin- and caveolae-mediated endocytosis, respectively. HeLa cells treated with UCNP-PEI without inhibitors were used as controls.