Electronic Supplementary Information

An upconversion nanoplatform for simultaneous photodynamic

therapy and Pt chemotherapy to combat cisplatin resistance

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Figure S1. Photographs of PAA-UCNPs and PEG/RB-Pt(IV)-UCNPs (100 μ g mL⁻¹) in aqueous solutions before and after centrifugation.



Figure S2. ¹H NMR spectrum of *c,c,t*-[Pt(NH₃)₂Cl₂(OCOCH₂CH₂NH₃)₂]²⁺·2CF₃COO⁻ (DMSO-*d*₆)



Figure S3. ¹³C NMR spectrum *c,c,t*-[Pt(NH₃)₂Cl₂(OCOCH₂CH₂NH₃)₂]²⁺·2CF₃COO⁻ (DMSO-*d*₆)



Figure S4. ¹⁹⁵Pt NMR spectrum of *c*,*c*,*t*-[Pt(NH₃)₂Cl₂(OCOCH₂CH₂NH₃)₂]²⁺·2CF₃COO⁻ (DMSO-*d*₆)



Figure S5. Mass spectrum of $\{c, c, t-[Pt(NH_3)_2Cl_2(OCOCH_2CH_2NH_2)_2]+H\}^+$.



Figure S6. Photoluminescent spectrum of PAA-UCNPs under 808 nm laser irradiation at 6 W/cm² (black) and UV-Vis spectrum of RB (red).



Figure S7. UV-vis spectra of RB-Pt(IV)-UCNPs with different RB loading amount (0, 0.2, 0.6, 1.2, 2.6, 4.0 wt% RB on UCNPs).



Figure S8. Photoluminescent spectra of PAA-UCNPs, Pt(IV)-UCNPs, and RB-Pt(IV)-UCNPs.



Figure S9. Release of RB and Pt from PEG/RB-Pt(IV)-UCNPs in PBS buffer (120 μ g mL⁻¹) at pH 7.4 at r.t.



Figure S10. Pt release profile of PEG/RB-Pt(IV)-UCNPs in PBS buffer (400 μ g mL⁻¹) in and without the presence of 2 mM ascorbic acid (AsA) at pH 7.4 at 37 °C. One batch of sample was irradiated with an 808 nm laser for 10 min (6 W/cm²) before the release experiment was carried out.



Figure S11. Whole cell uptake of cDDP (11.4 μ M) and PEG/RB-Pt(IV)-UCNPs (80 μ g mL⁻¹) in A2780cisR cells (n=3).



Figure S12. Cellular reactive oxygen species (ROS) of PEG/RB-Pt(IV)-UCNPs (80 μ g mL⁻¹) with 808 nm laser irradiation (6 W/cm², 10 min) in A2780cisR cells. The PBS treated and PEG/RB-Pt(IV)-UCNPs (80 μ g mL⁻¹) without irradiation were also measured to compare. The H₂O₂ (3 mM) treated for 15 min was set as a positive control. Scale bar, 250 μ m.



Scheme S1. Synthetic route of PEG/RB-HA-UCNPs for photodynamic therapy alone.