

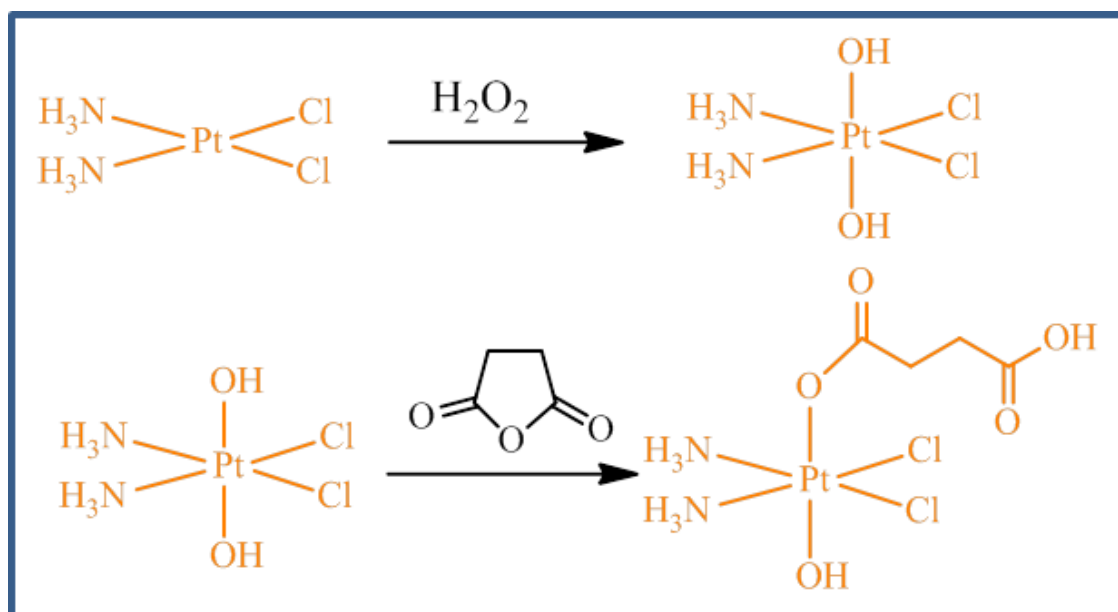
Supporting Information

Dextran–platinum(IV) conjugate as reduction-responsive carrier for triggered drug release

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Synthesis of *c,c,t*-[Pt(NH₃)₂Cl₂(OH)₂]

Briefly, H₂O₂ (5 mL) was added to a suspension of cisplatin (1 g) in H₂O (5 mL), and the mixture was stirred at room temperature in dark for 4 h. Then the light yellow precipitate was collected and washed with water and acetone for several times, and dried (Scheme S1). ¹H NMR: (300 MHz, DMSO-d₆, 25 °C): 5.2-5.8 (6H, m, NH₃); IR: cm⁻¹ 3463 (br, OH), 538 (sh, Pt-OH). ¹H NMR: (300 MHz, d₆-DMSO, 25 °C): δH 2.33-2.39 (4H, m, -OCH₂CH₂C-), 5.12-5.71 (6H, m, NH₃) (Fig. 2A, Fig. S1).



Scheme S1 Synthesis of *c,c,t*-[Pt(NH₃)₂Cl₂(OH)₂] and Pt(IV).

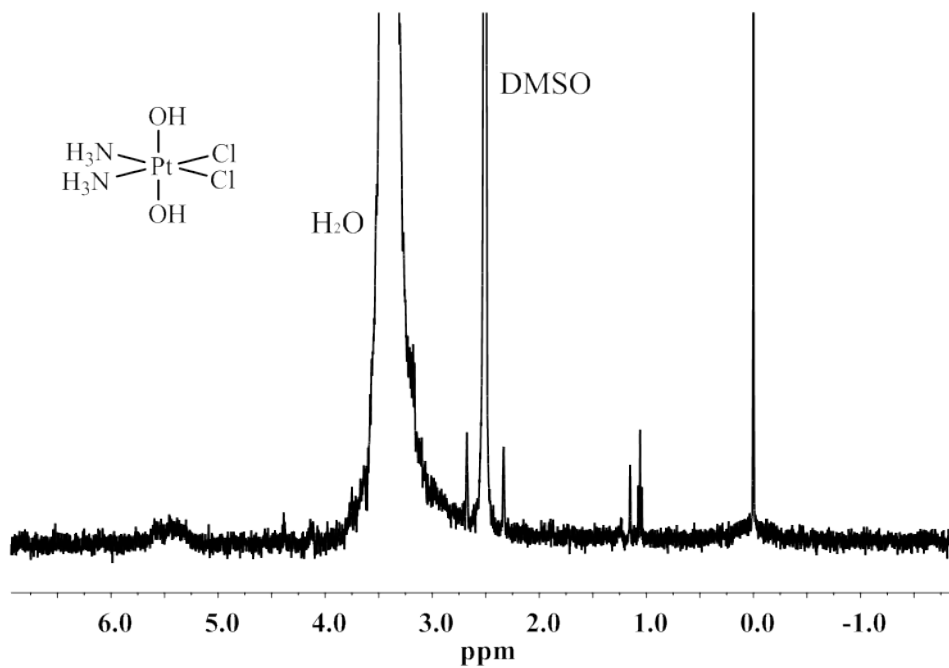


Fig. S1 The ¹H NMR spectra of c,c,t -[Pt(NH₃)₂Cl₂(OH)₂].

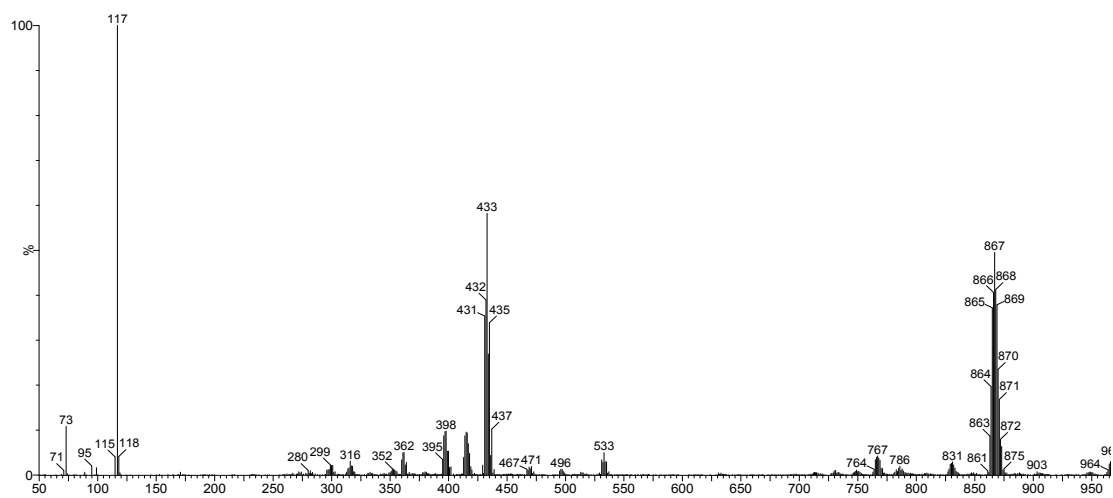


Fig. S2 ESI-MS spectrum of Pt(IV).

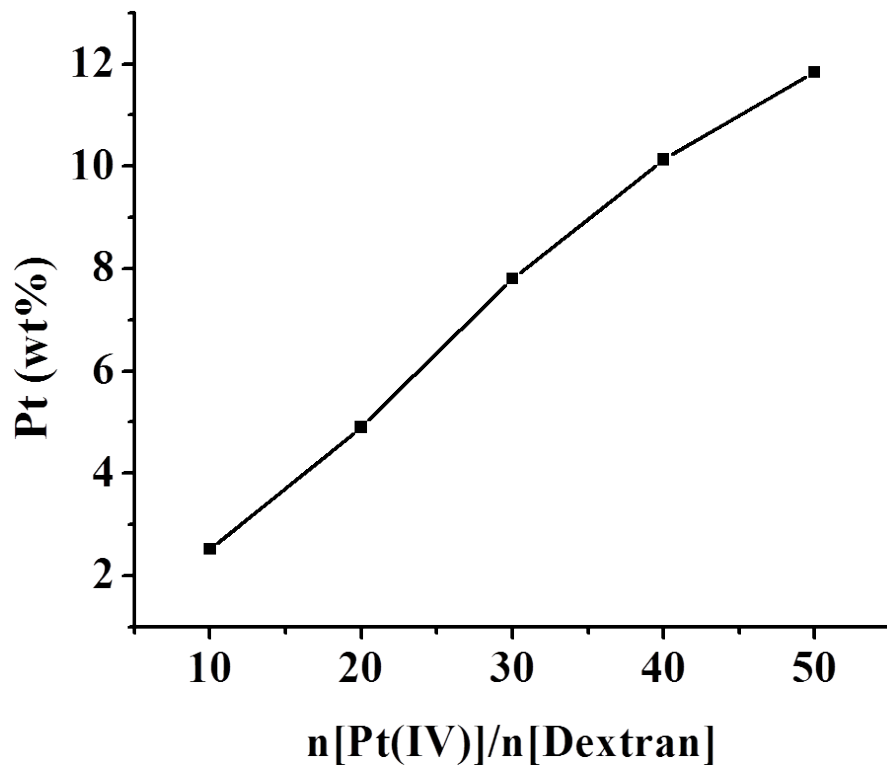


Fig. S3 Platinum content of the synthesized Dextran–Pt(IV) conjugate as a function of the molar ratio of Pt(IV) to dextran.

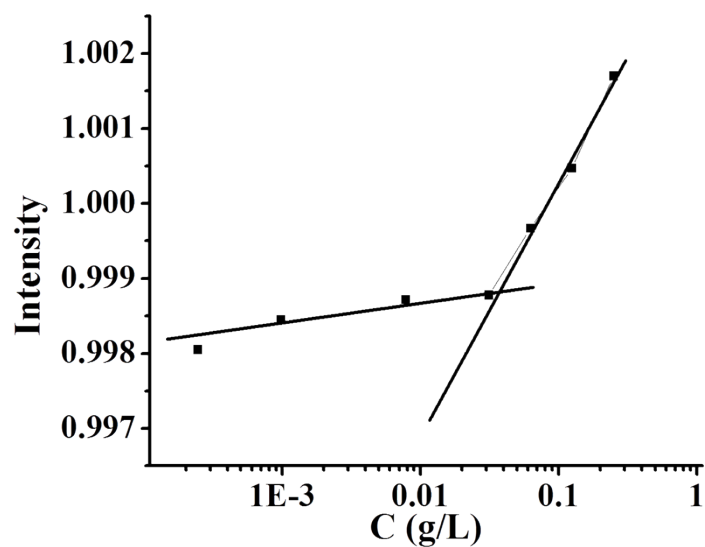


Fig. S4 Intensity of pyrene excitation spectra vs. the logarithm of the concentration of Dextran–Pt(IV) conjugate in water.

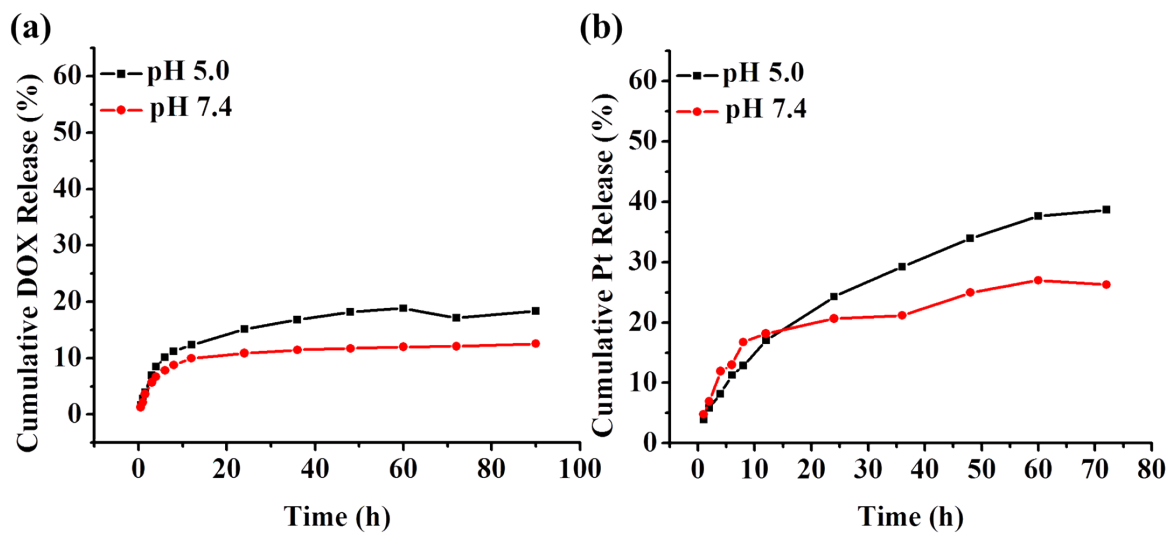


Fig. S5 *In vitro* drug release profiles of DOX-loaded Dextran–Pt(IV) in PBS (pH 7.4) and acetate buffer solution (pH 5.0): (a) DOX release profiles; (b) Pt release profiles.