

Electronic Supplementary Information

Multifunctional magnetic calcium phosphate nanoparticles for targeted platinum delivery

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Figure S1

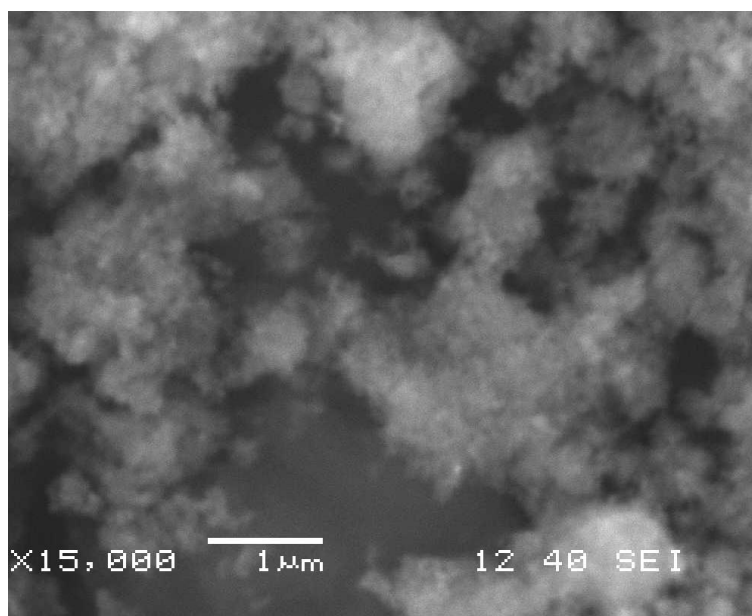


Figure S1 SEM image of CoFe₂O₄@ACP-CDDP-FA-RITC nanoparticles

Figure S2

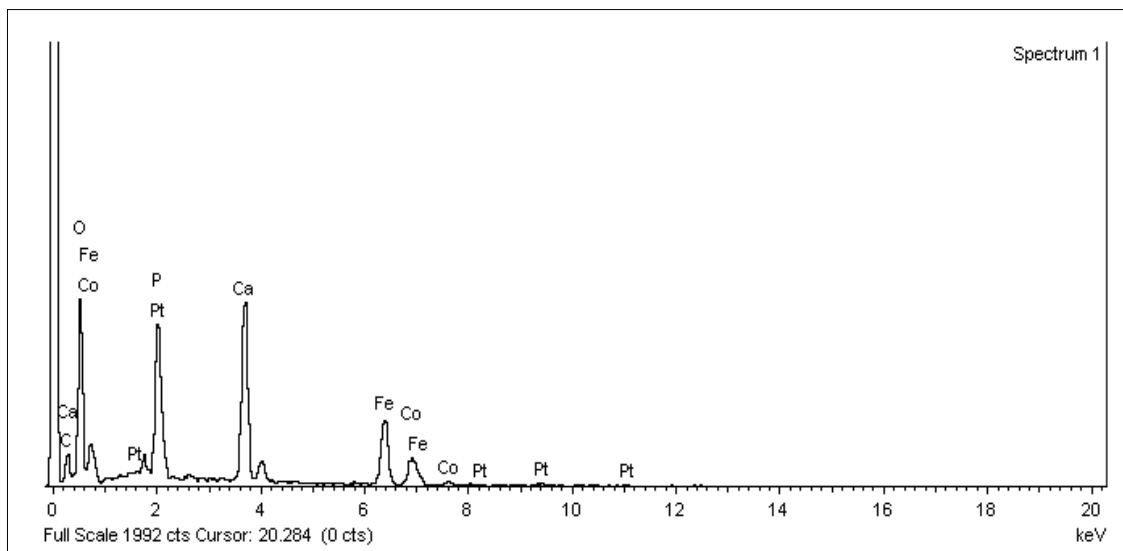


Figure S2 EDX spectrum of $\text{CoFe}_2\text{O}_4@ACP\text{-CDDP-FA-RITC}$ nanoparticles

Figure S3

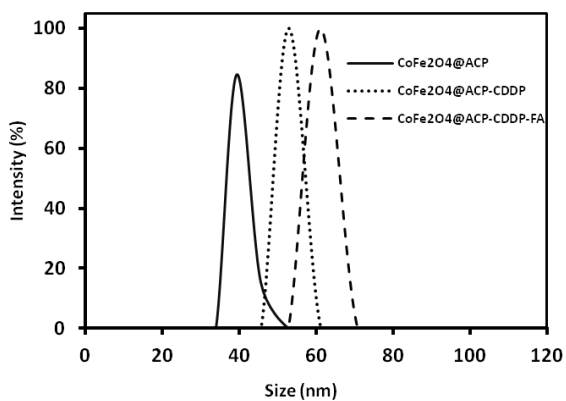


Figure S3 Variation of hydrodynamic size after each step of modification

Figure S4

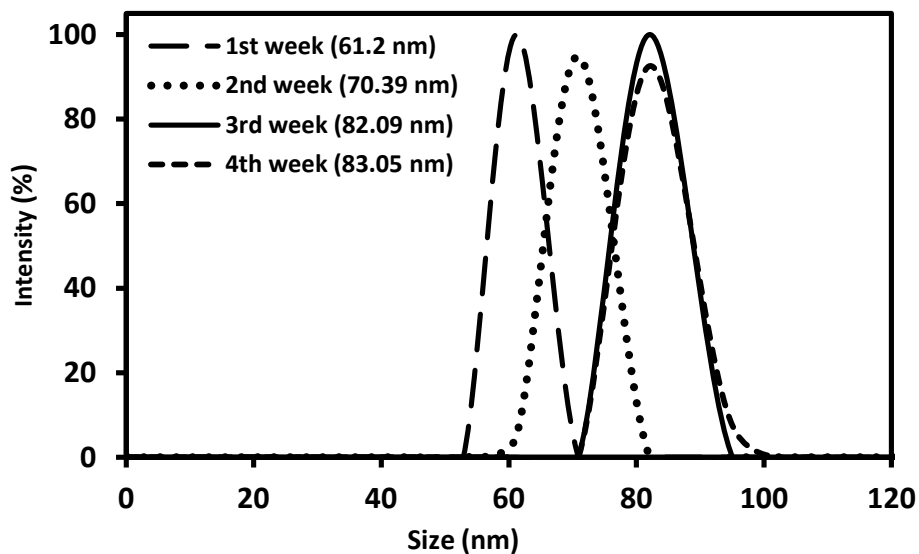


Figure S4 Change in hydrodynamic size of CoFe₂O₄@ACP-CDDP-FA-RITC with respect to time

Figure S5

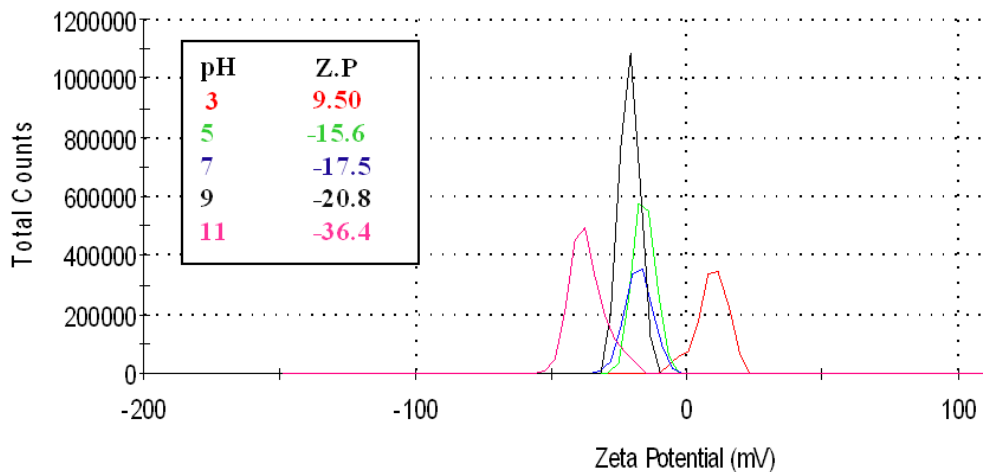


Figure S5 Zeta potential variation of CoFe₂O₄@ACP-CDDP-FA with respect to pH

Figure S6

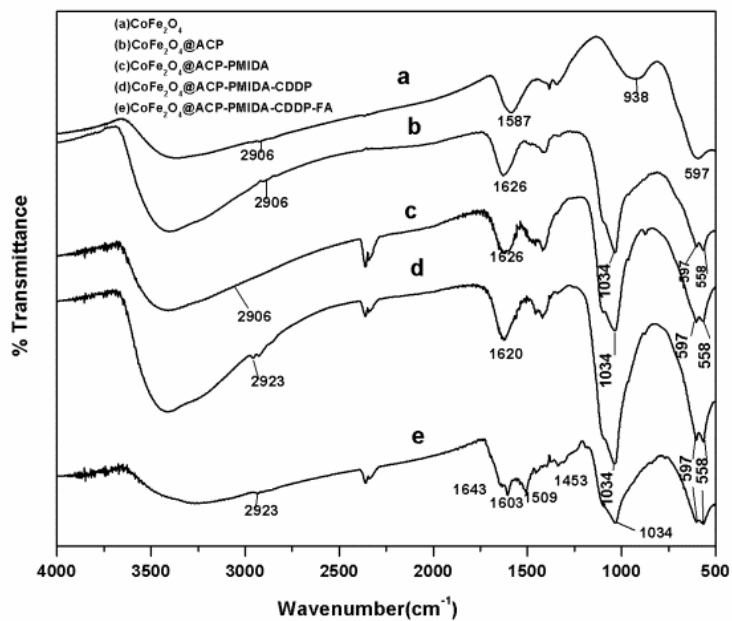


Figure S6 FTIR spectra after each step of conjugation

Figure S7

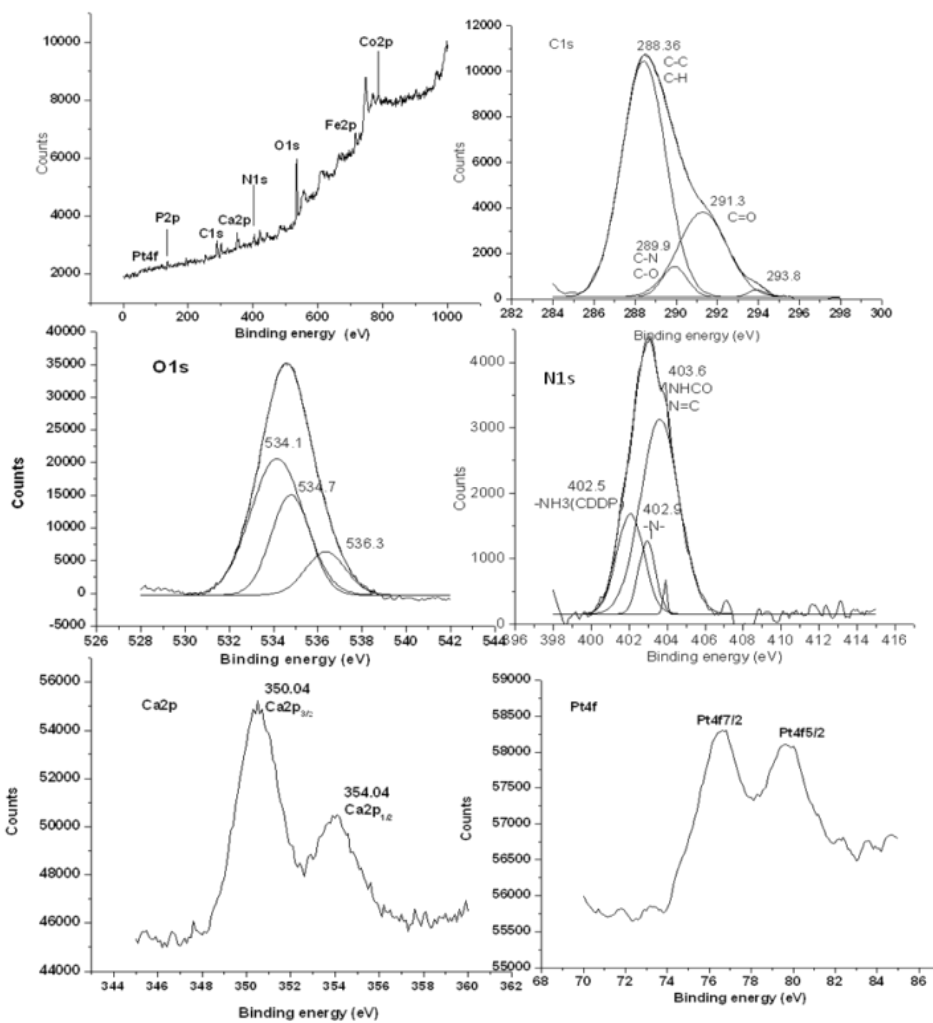


Figure S7 XPS survey spectrum of CoFe₂O₄@ACP-CDDP-FA. High resolution scans of (b) C1s, (c) O1s, (d) N1s, (e) Ca2p, (f) Pt4f

Figure S8

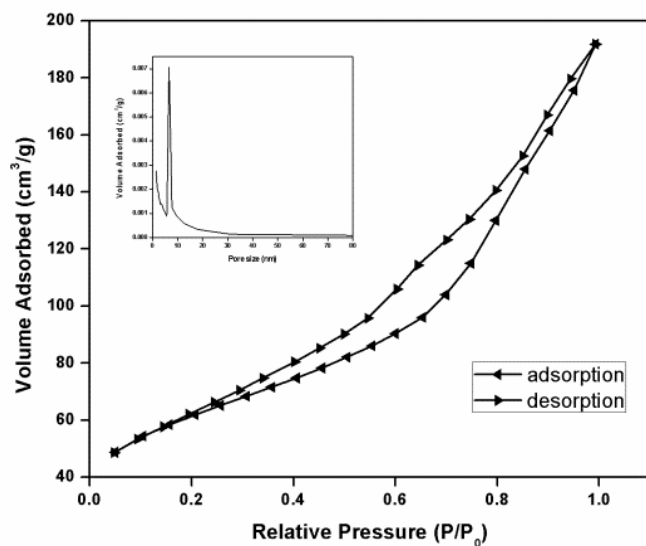


Figure S8 Nitrogen adsorption-desorption isotherm and the corresponding pore size distribution (inset) of as synthesized CoFe₂O₄ nanoparticles

Figure S9

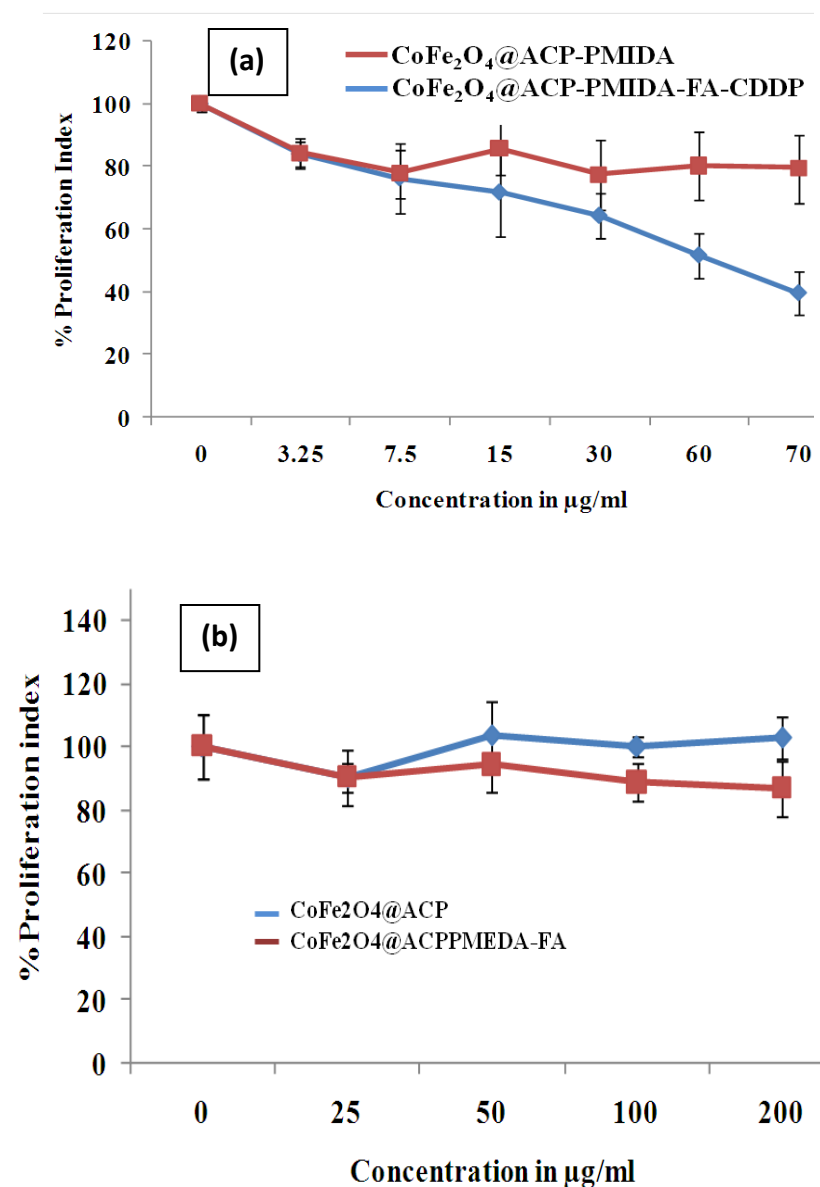


Figure S9 Cytotoxicity assay of nanoparticles on (a) HeLa and (b) L929 cells. Cells were treated with different concentrations of nanoparticles for 72 h and cell viability was measured by MTT assay

Figure S10

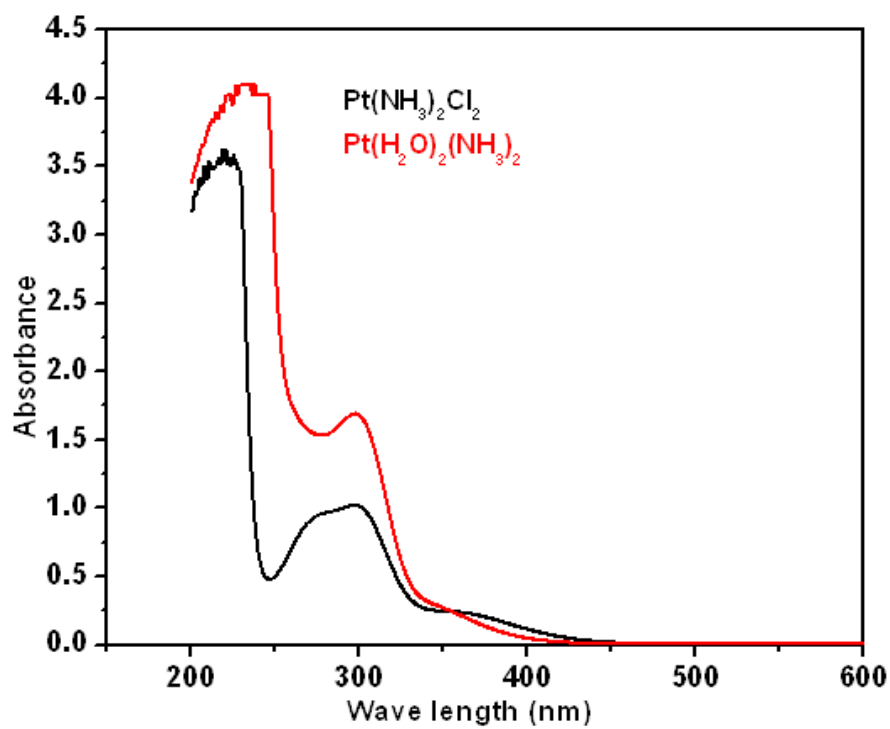


Figure S10 Change in UV-Visible spectrum of $\text{Pt}(\text{NH}_3)_2\text{Cl}_2$ after reacting with 2 equivalents of AgNO_3

Figure S11

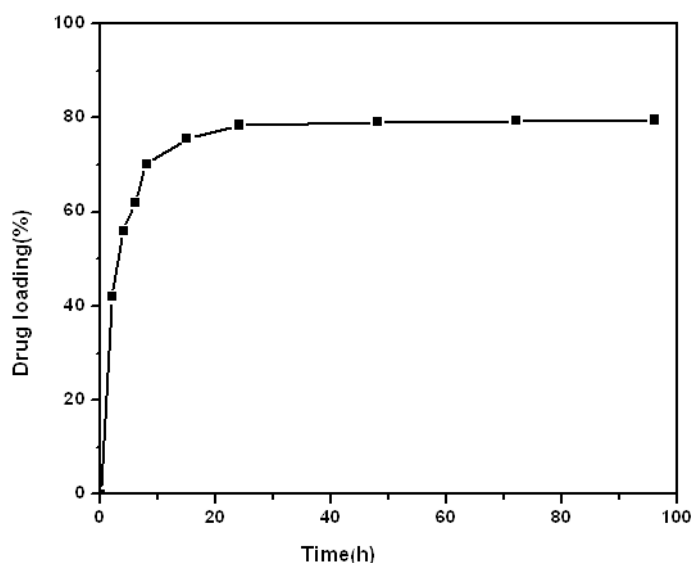


Figure S11 CDDP loading (%) on -COOH functionalized CoFe₂O₄ @ ACP nanoparticles

Calculation of drug loading capacity

The drug loading capacity was calculated as per our previously reported paper [1]. First, COOH functionalized particles were conjugated with Pt(H₂O)₂Cl₂ as described in the experimental section and then particles were separated from the aqueous suspension using magnetic separator (Invitrogen). The obtained drug-loaded CoFe₂O₄@ACP nanoparticles were incubated at 60 °C in vacuum overnight and were weighted. Drug concentration in supernatant was analyzed by the ultraviolet absorption ($\lambda_{\text{max}} = 295 \text{ nm}$), with reference to a calibration curve on a UV-Vis spectrophotometer. The measurements were performed in triplicate. Drug-loading content and encapsulation efficiency were obtained by eqs 1 and 2, respectively [2].

$$\begin{aligned} \text{Drug-loading content (\%)} &= \frac{\text{Weight of the drug in nanoparticles}}{\text{weight of the nanoparticles}} \times 100 \\ &= \frac{0.013}{0.120} \times 100 = 10.8\% \end{aligned}$$

$$\begin{aligned} \text{Encapsulation efficiency (\%)} &= \frac{\text{Weight of the drug in nanoparticles}}{\text{Weight of the feeding drug}} \times 100 \\ &= \frac{0.013}{0.017} \times 100 = 76.4\% \end{aligned}$$