Magnetic/NIR-responsive drug carrier, multicolor cell imaging, and enhanced photothermal therapy of gold capped magnetite-fluorescent carbon hybrid nanoparticles

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Figure S1. Synthetic process of multifunctional $Fe_3O_4@PC-CDs-Au$ NPs. Stage I: formation of $Fe_3O_4@PC-CDs-Ag$ NPs by the loading and reduction of Ag^+ ; Stage II: formation of $Fe_3O_4@PC-CDs-Au$ NPs by the galvanic replacement reaction between the Au precursor (HAuCl₄) and Ag nanocrystals.

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Figure S2. Typical XRD pattern of the as-obtained Fe₃O₄@PC-CDs NPs.



Figure S3. FT-IR spectra of the Fe₃O₄@PC-CDs and Fe₃O₄@PC-CDs-Ag hybrid NPs.



Figure S4. (A) TEM image of the as-obtained Fe₃O₄@PC-CDs-Ag NPs ;(B) EDAX of the single Fe₃O₄@PC-CDs-Ag NP.



Figure S5. (A) PL spectra of the Fe₃O₄@PC-CDs-Au NPs at different excitation wavelengths. (B) Time dependent PL intensity variation (7.5%) of the Fe₃O₄@PC-CDs-Au NPs under a 2h continuous exposure to the excitation UV light of 365 nm.





 $\label{eq:Figure S7. Z-Scanning confocal fluorescence transmission images of mouse melanoma cells B16F10 incubated with $$Fe_3O_4@PC-CDs-Au NPs.$$$



Figure S8. The pore size distribution of the as-obtained Fe₃O₄@PC-CDs-Au NPs.



Figure S9. Hyperthermia assay on aqueous dispersion of $Fe_3O_4@PC-CDs-Au$ NPs (0.01 g/L) and water (control) under an alternating magnetic field.



Figure S10. (A) Releasing profiles of DOX from the Fe₃O₄@PC-CDs-Au NPs at pH=7.4 under different temperatures of 27 °C, 37 °C, and 42 °C, respectively. (B) Releasing profiles of DOX from the Fe₃O₄@PC-CDs-Au NPs in buffer solutions of different pH of 5.0 and 7.4, respectively, at 37 °C.



Figure S11. In vitro cell viability in the control culture medium without/with 1.5 W/cm² NIR irradiation for 5 min.

m².