

Supporting Information

Multifunctional Fe₃O₄@Polydopamine Core-Shell Nanocomposites for Intracellular mRNA Detection and Imaging-Guided Photothermal Therapy

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Table S1. The sequences employed in this work.

Name	Sequence
FAM-hpDNA	5'- <u>CCGGGTTTGGTGAAGCTAACGTTGAGGACCCGG</u> -FAM-3'
c-myc target	5'-CCUCAACGUUAGCUUCACCAA-3'
non-complementary target	5'-CAAGGAGCUGGAAGGCUGGGA-3'
Cy3-hpDNA	5'- <u>CCGGGTGCGAGTGTCTTTGGCATACTTACCCGG</u> -Cy3-3'

Underline denotes base pairs in the stem.

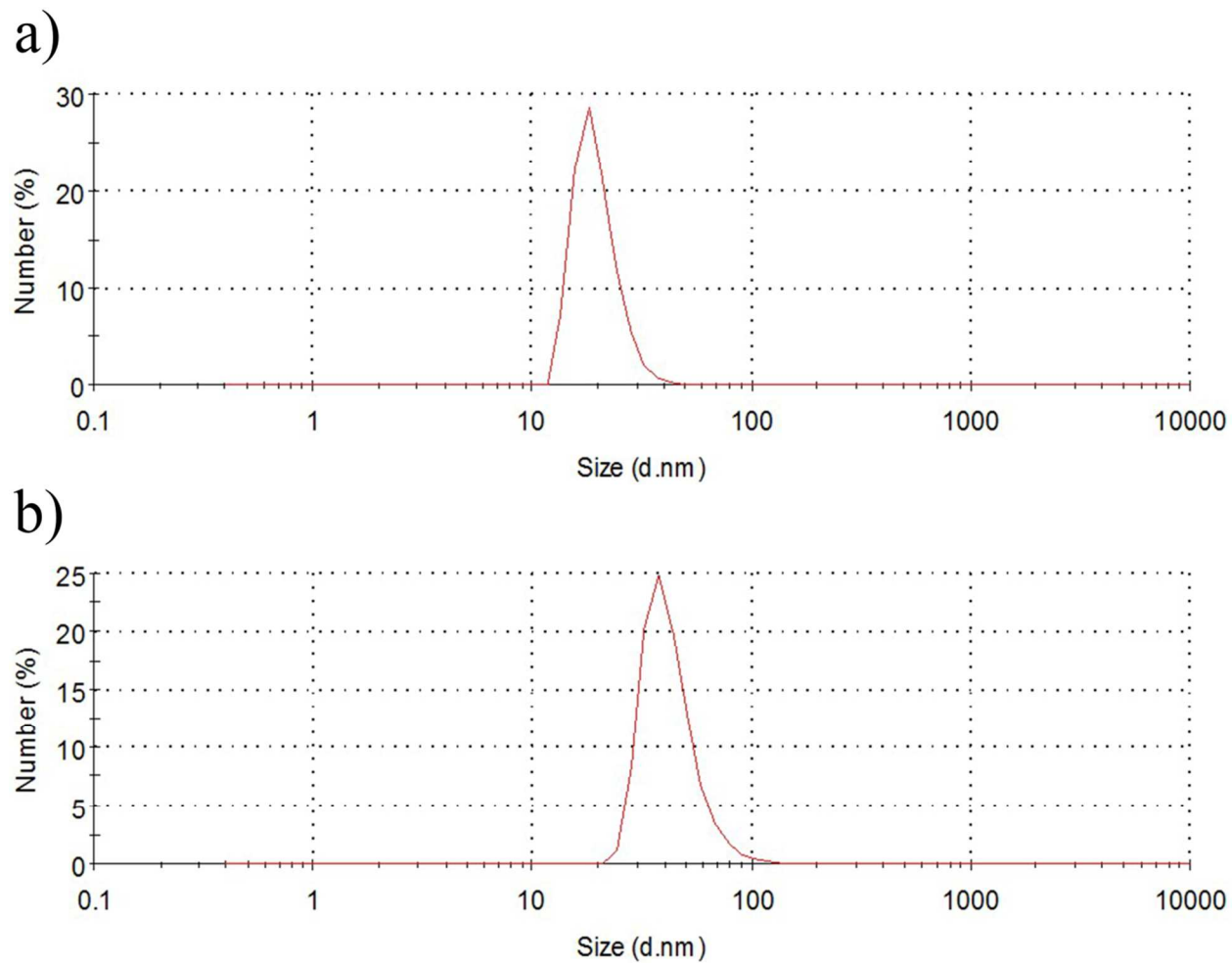


Figure S1. Dynamic light scattering (DLS) data of (a) Fe_3O_4 NPs and (b) Fe_3O_4 @PDA NCs in water.

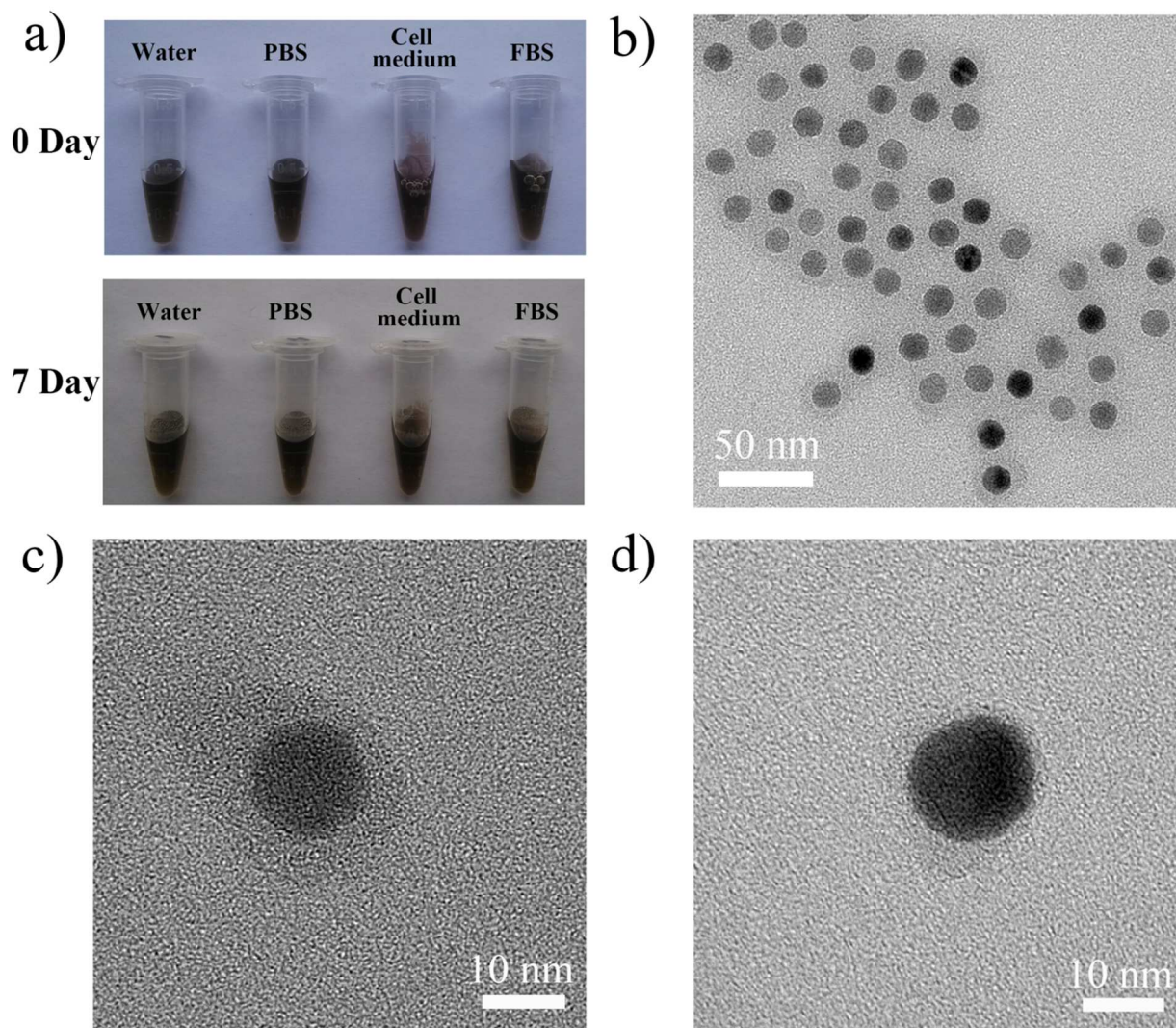


Figure S2. (a) Photos of Fe₃O₄@PDA NCs in different solutions including water, PBS, RPMI-1640 with 10% fetal bovine serum (FBS) cell culture medium, and FBS. The Fe₃O₄@PDA NCs were stable in those solutions for at least one week without showing obvious aggregation. (b,c,d) TEM images of the Fe₃O₄@PDA NCs prepared with different polymerization time: 4 h (b,c) and 1 h (d).

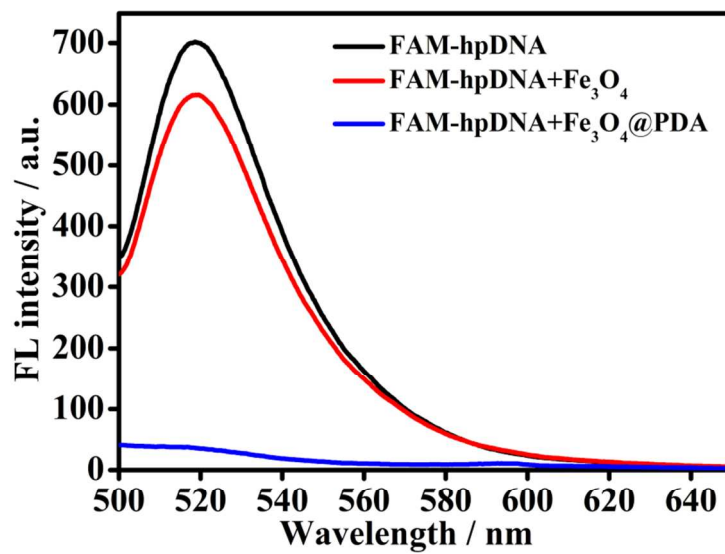


Figure S3. Fluorescence quenching of 50 nM FAM-hpDNA in the presence of Fe₃O₄ NPs or Fe₃O₄@PDA NCs. Excitation: 480 nm, emission: 520 nm.

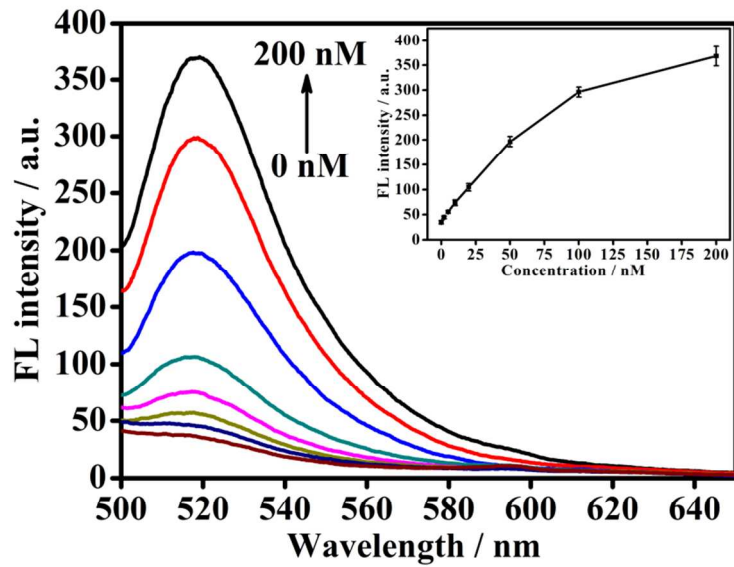


Figure S4. Fluorescence emission spectra of Fe₃O₄@PDA-DNA nanoprobe treated with various concentrations of RNA target (0, 2, 5, 10, 20, 50, 100 and 200 nM). Excitation: 480 nm, emission: 520 nm.

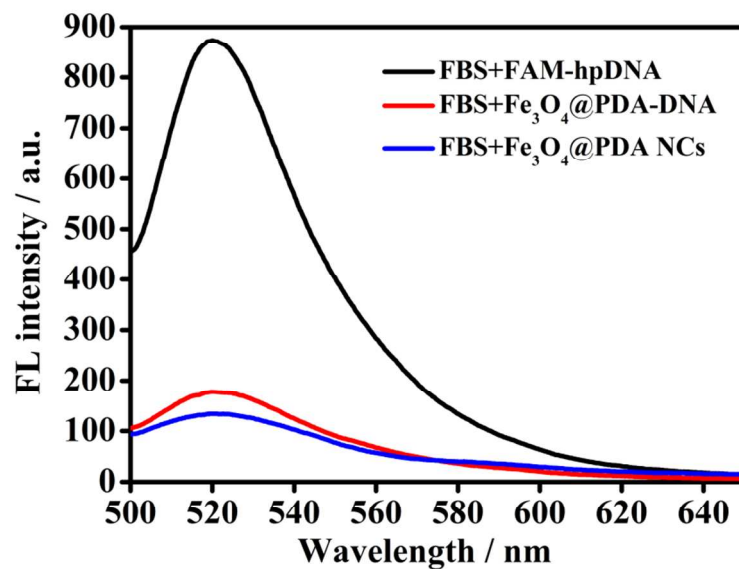


Figure S5. Fluorescence emission spectra of 50 nM FAM-hpDNA or Fe₃O₄@PDA-DNA nanoprobe (containing 50 nM FAM-hpDNA) dispersed in FBS. Excitation: 480 nm, emission: 520 nm.

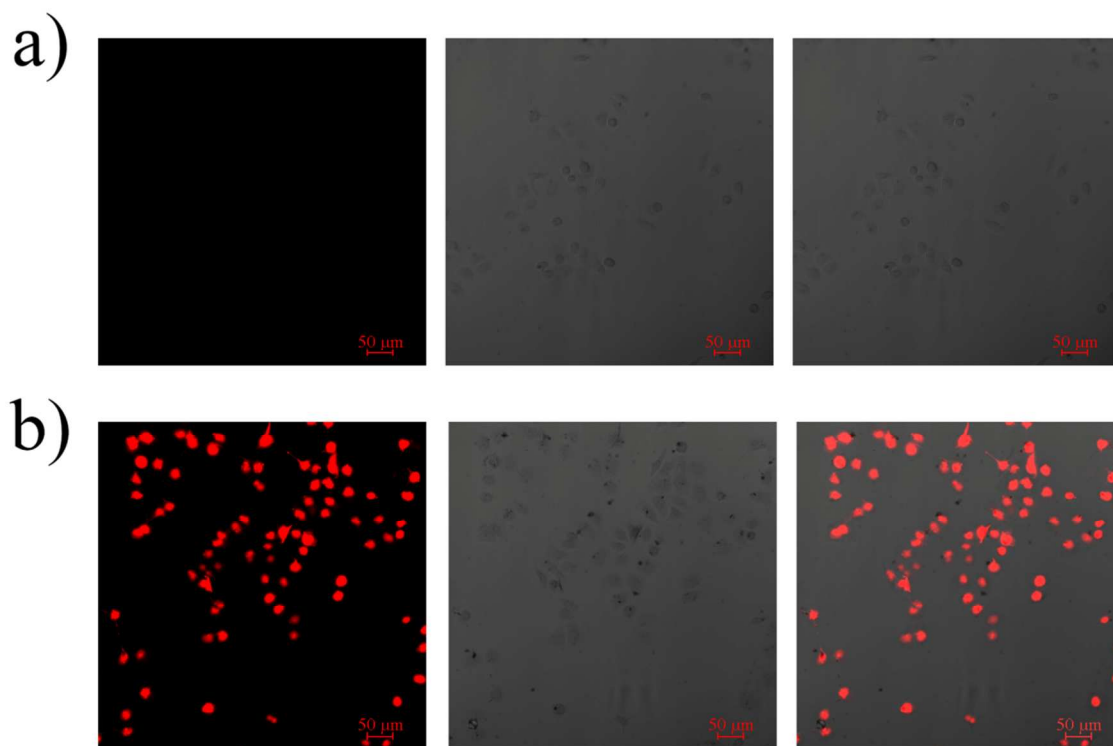


Figure S6. Propidium iodide (PI) stained CLSM images of MCF-7 cells incubated with 0.1 mg mL^{-1} Fe_3O_4 NPs (a) or $\text{Fe}_3\text{O}_4@PDA$ NCs (b) after laser irradiation for 5 min. Left panels are PI fluorescence corresponding to dead cells, center panels are bright field image of cells, and right panels are the overlay of PI fluorescence and the bright field image. Scale bars are $50 \mu\text{m}$.

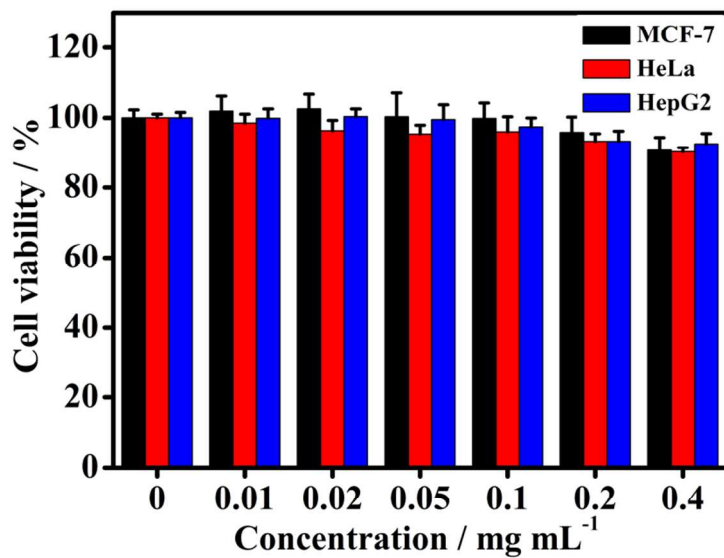


Figure S7. Cell viability of MCF-7, HeLa, and HepG2 cells incubated with different concentrations of Fe₃O₄@PDA NCs for 24 h.