

Surface Modified Multifunctional ZnFe₂O₄ Nanoparticles for Hydrophobic and Hydrophilic Anti-Cancer Drug Molecules Loading

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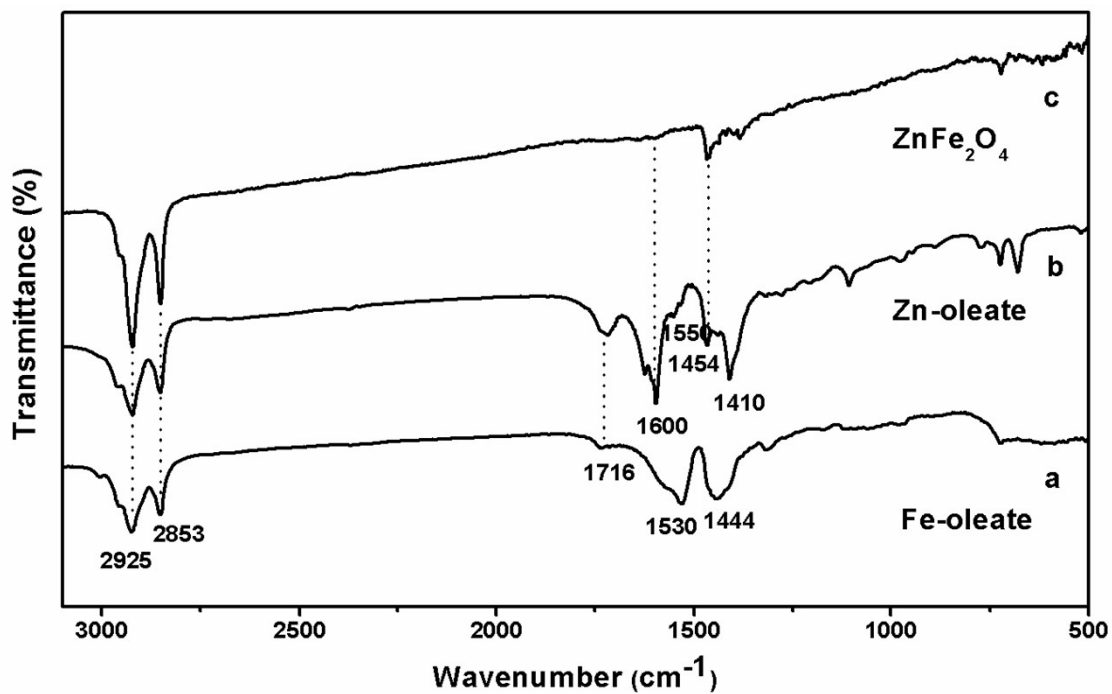


Fig. S1: FT-IR spectra of (a) Fe-Oleate (b) Zn-Oleate (c) as prepared ZnFe_2O_4 nanoparticles

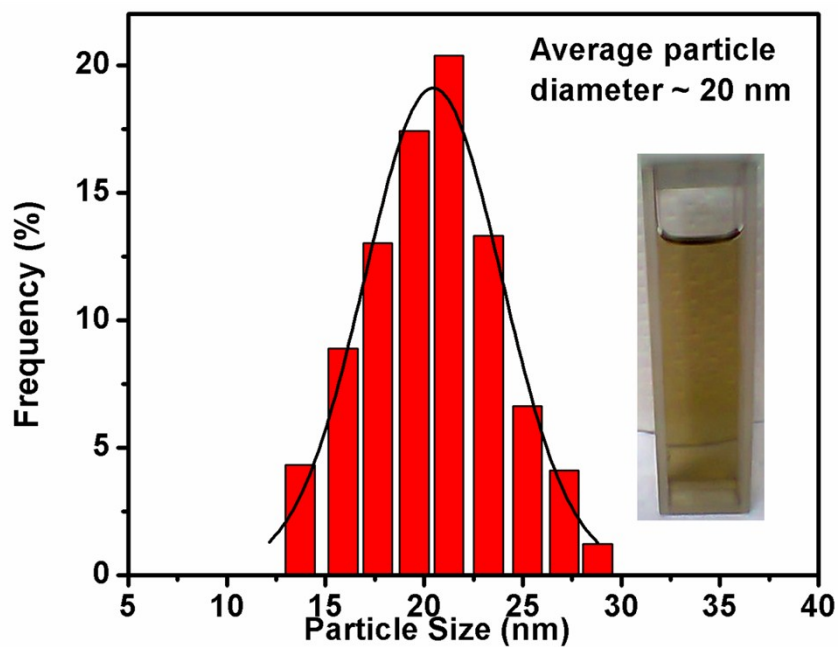


Fig. S2: The measured particle size distribution of (a) chloroform dispersed ZnFe_2O_4 nanoparticles (inset) and the average particles size is ~20 nm.

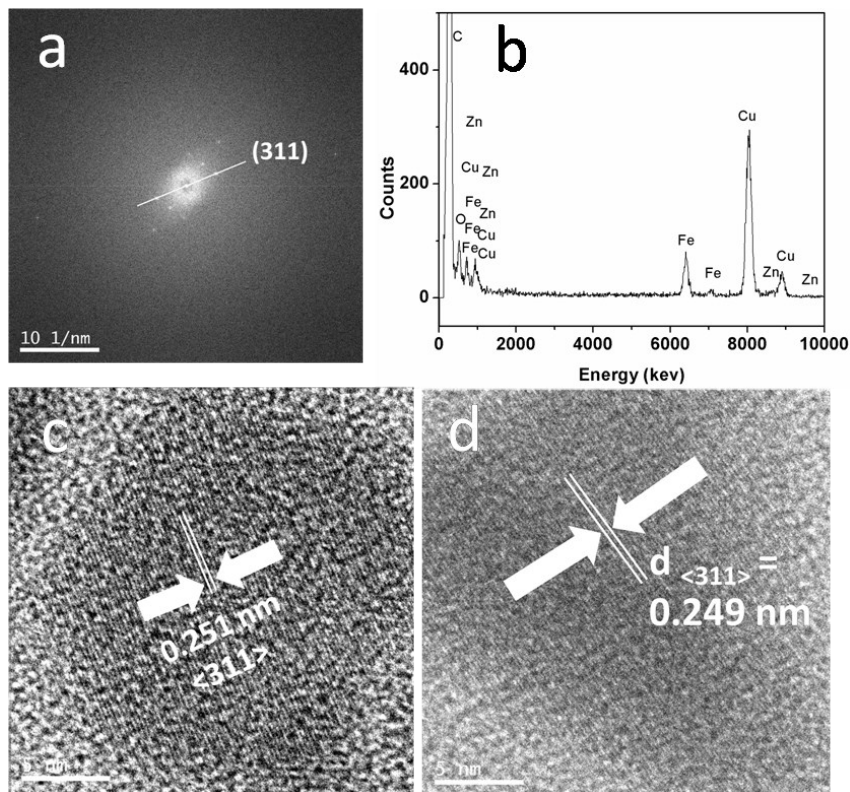


Fig. S3: (a) Fast Fourier transformation (FFT) pattern clearly indicates the mono crystalline nature of the particles which have 311 planes of ZnFe_2O_4 (b) Energy-dispersive X-ray spectroscopy (EDS) shows the presence of Zn and Fe elements in the sample (c) and (d) are HRTEM images of ZnFe_2O_4 nanoparticles.

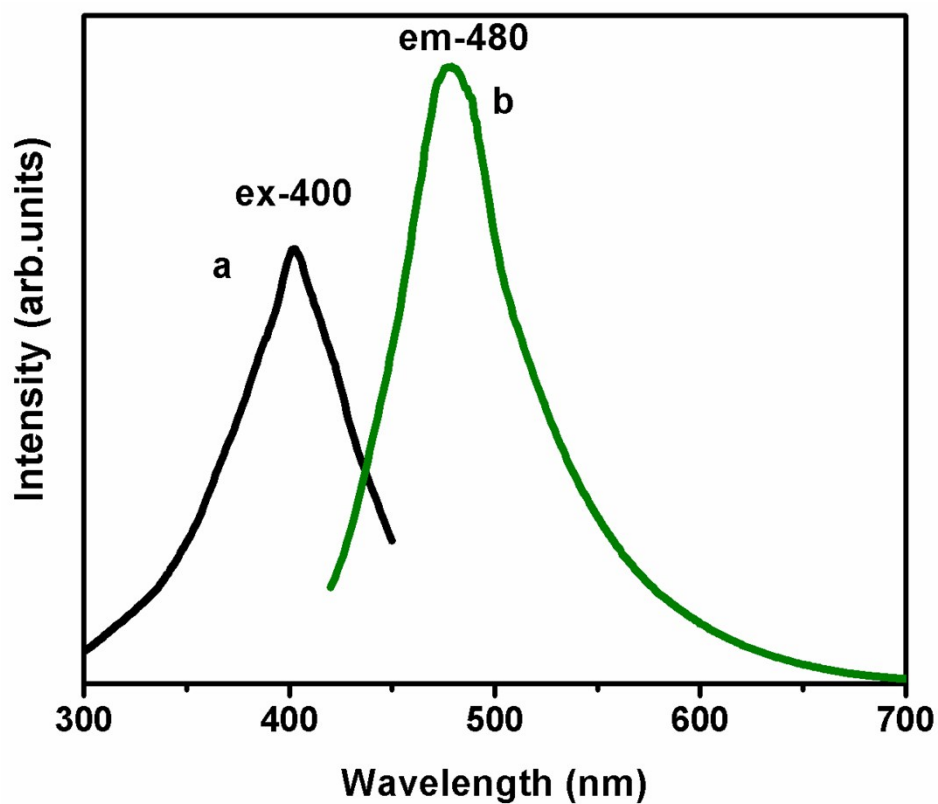


Fig. S4: (a) The excitation spectrum at $\lambda_{\text{emission}}$ 480 nm and (b) Emission spectrum at $\lambda_{\text{excitation}}$ 400 nm of ZnFe_2O_4 nanoparticles.

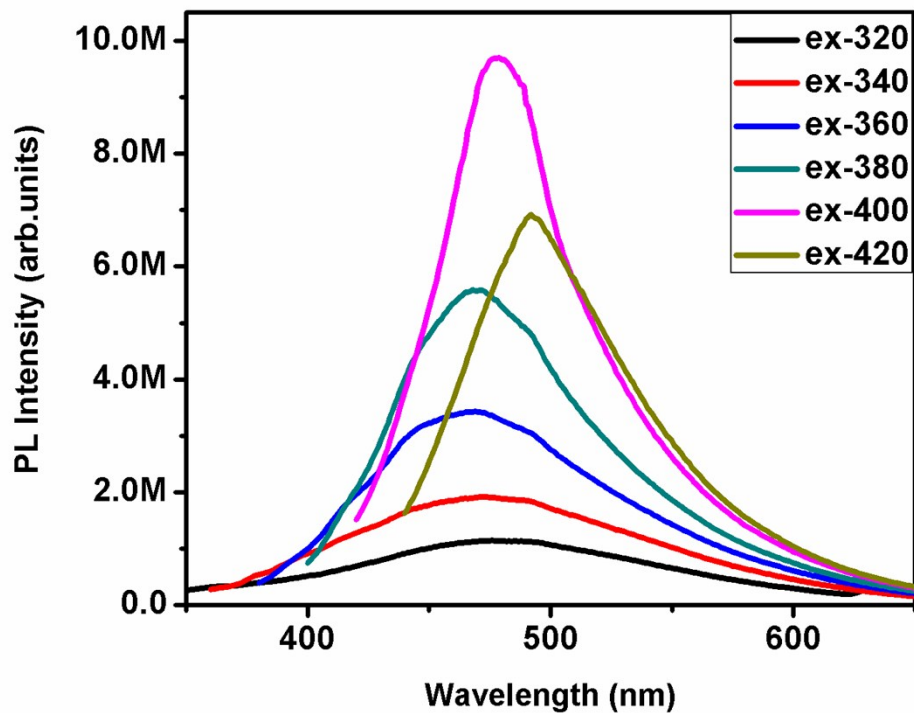


Fig. S5: The photoluminescence spectra of chloroform dispersed ZnFe₂O₄ nanoparticles. The solution was excited at different excitation wavelengths, $\lambda_{\text{excitation}}$ - 320, 340, 360, 380, 400 and 420 nm and emission peaks were centered at 470-480 nm. The highest intense peak was $\lambda_{\text{emission}}$ -480 nm corresponding to the $\lambda_{\text{excitation}}$ -400 nm

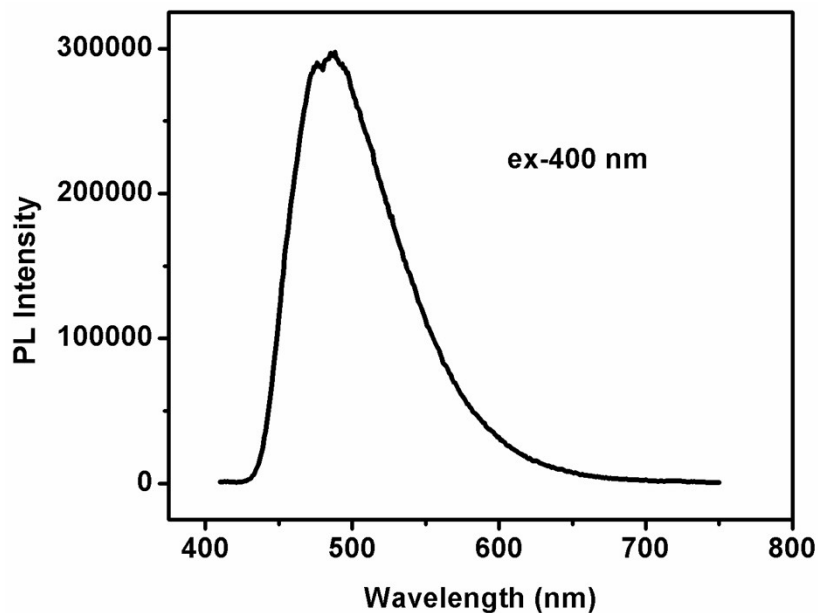


Fig. S6: Fluorescamine test performed to confirm the presence of primary amines groups on hydrophilic nanoparticle. For the test 200 μl of hydrophilic ZnFe_2O_4 solution and 100 μl of carbonate buffer solution were mixed together and fluorescamine was added the solution. The emission of the solution was carried upon excitation wavelength of 400 nm.

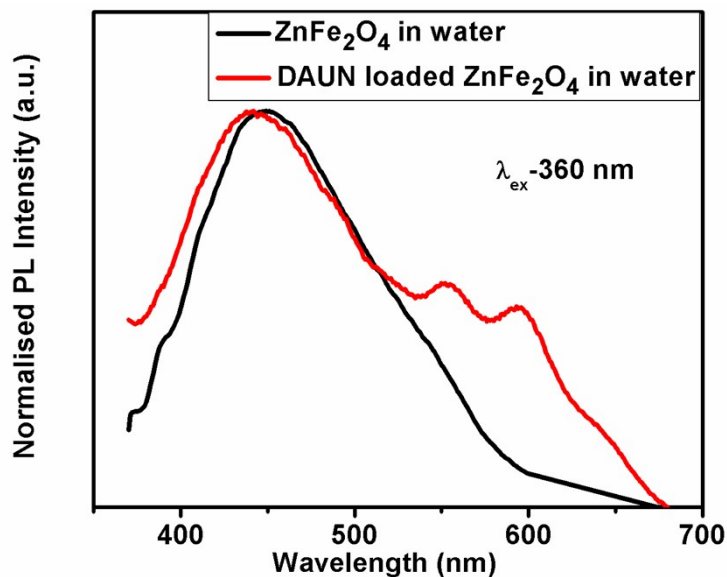


Fig. S7: The photoluminescence spectra of water dispersed ZnFe_2O_4 and DAUN loaded ZnFe_2O_4 nanoparticles on excitation at 360 nm.

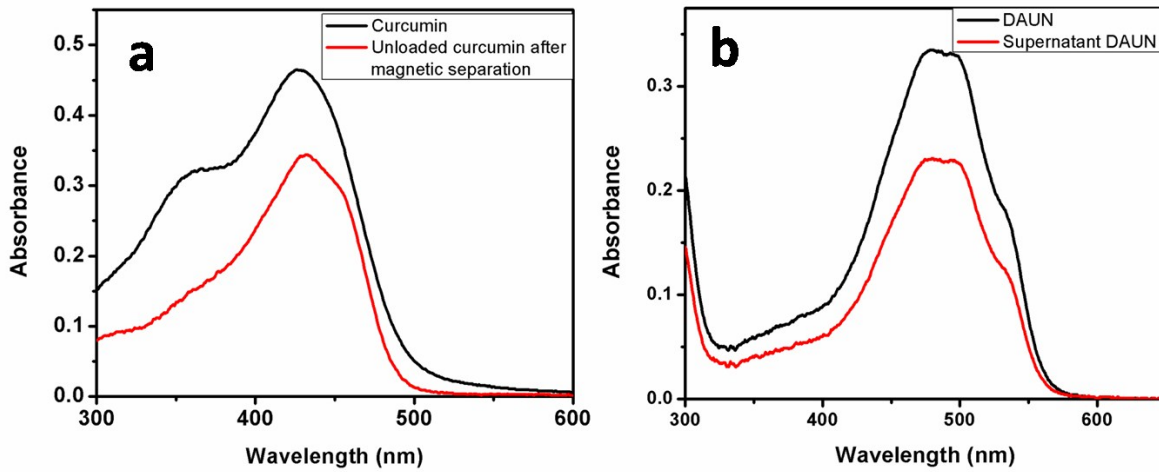


Fig. S8: UV-visible spectra for (a) Curcumin loading and (b) DAUN loading on ZnFe_2O_4 nanoparticles

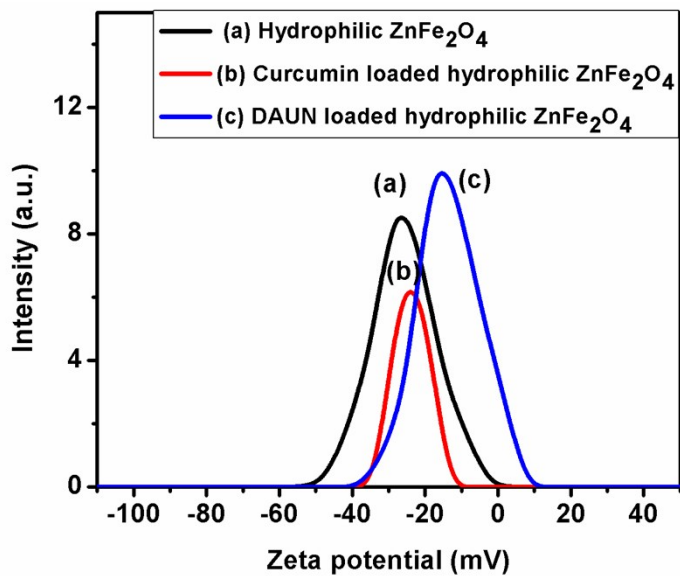


Fig. S9: Zeta potential measurement of hydrophilic and drug loaded ZnFe_2O_4 nanoparticle

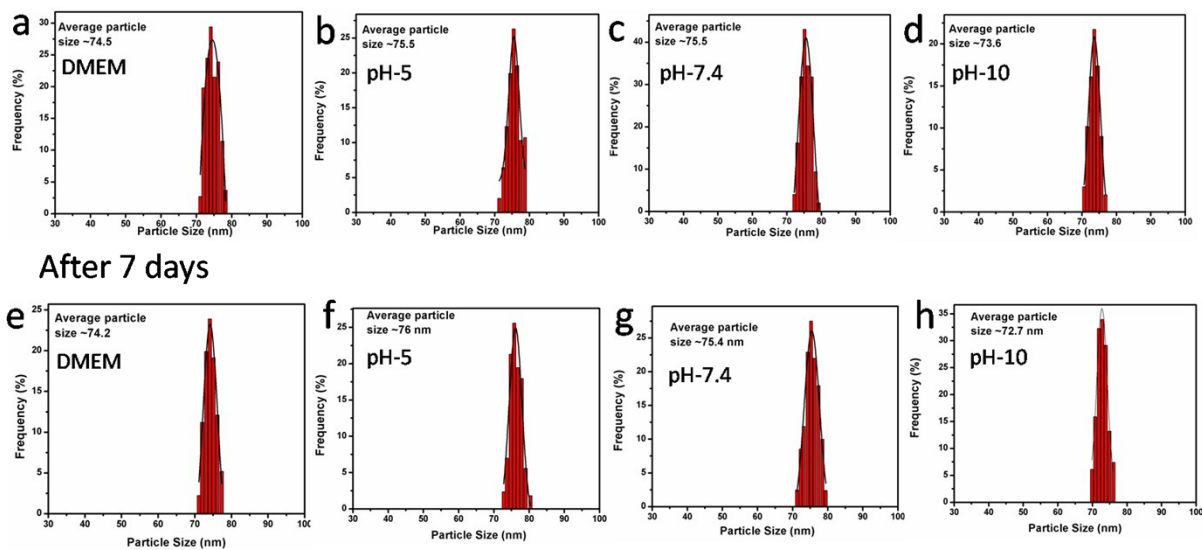


Fig. S10: Hydrodynamic particle size of ZnFe_2O_4 nanoparticles in DMEM and different pH-buffer solutions.

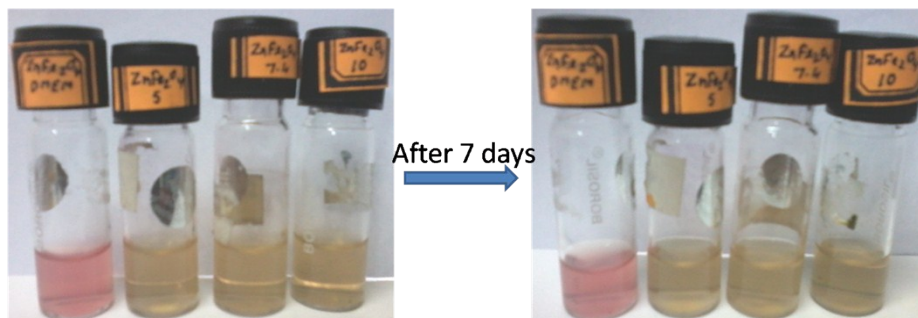


Fig. S11: Colloidal stability of ZnFe_2O_4 nanoparticles in DMEM and different pH-buffer solutions. Digital images show optically clear solutions.

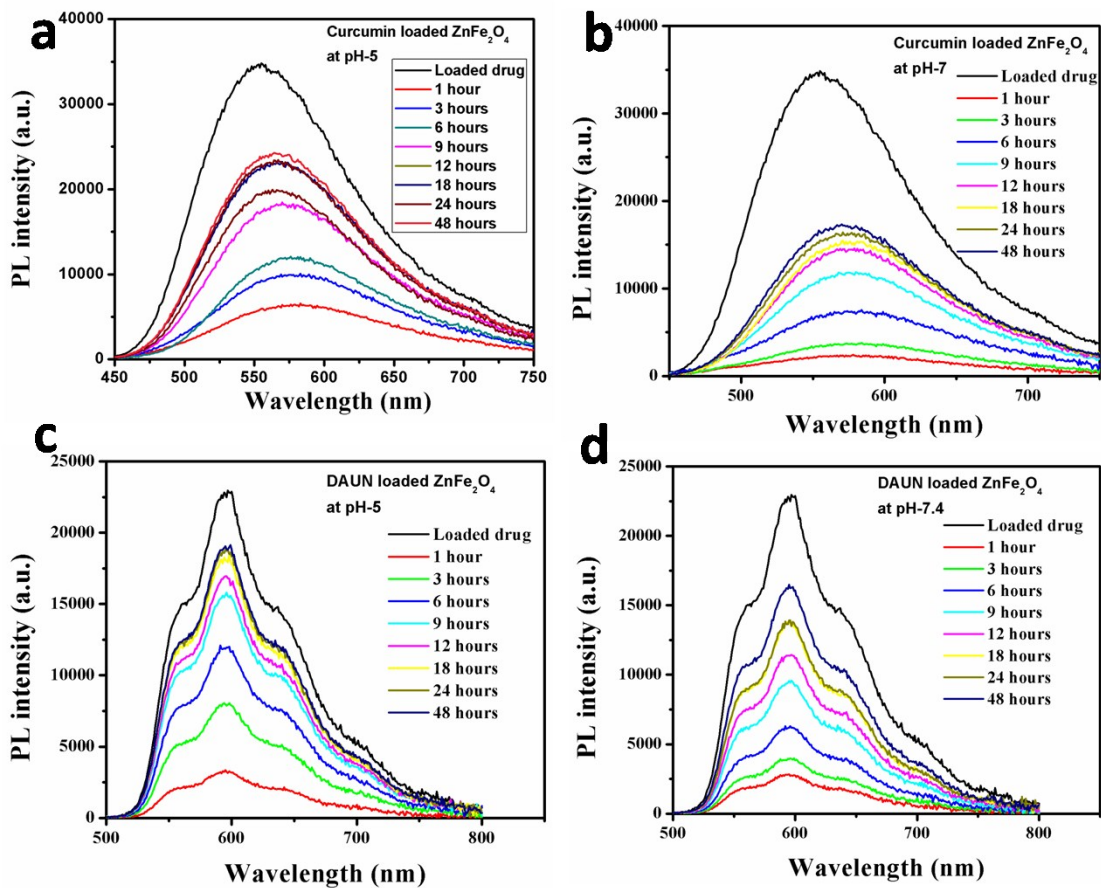


Fig. S12: Emission spectra depicting the drug release from curcumin loaded $ZnFe_2O_4$ nanoparticles a) at pH ~5 b) at pH ~7.4 and DAUN loaded $ZnFe_2O_4$ nanoparticles c) at pH ~5 and d) at pH ~7.4

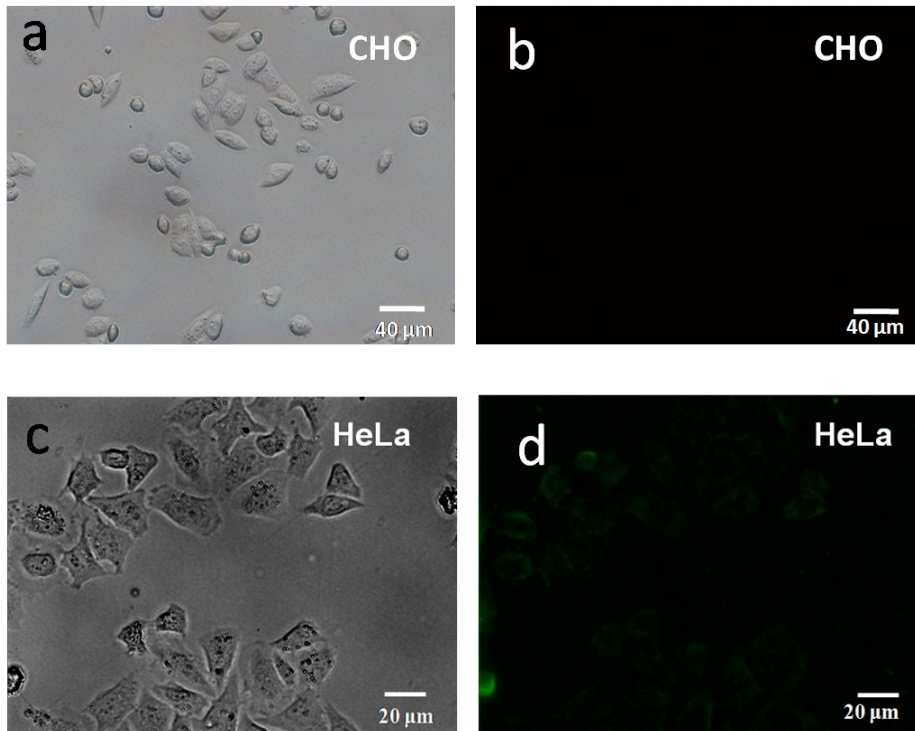
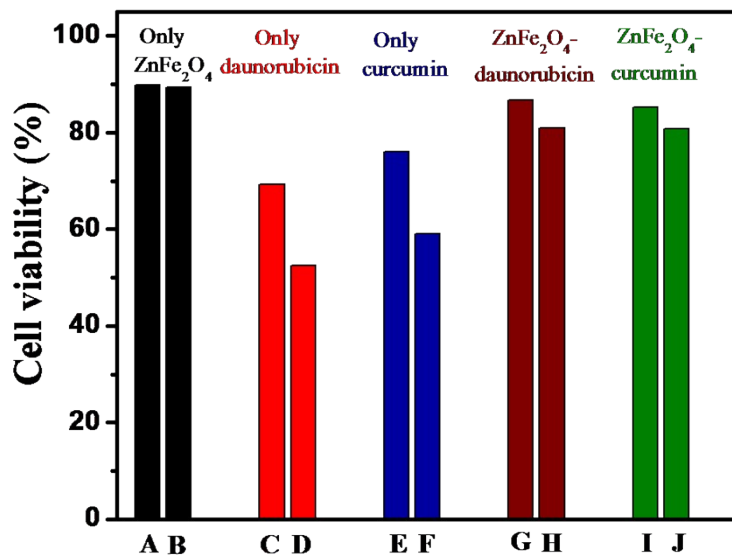


Fig. S13 The bright field (a,c) and fluorescence (b,d) imaging of CHO and HeLa cells respectively.



	A	B	C	D	E	F	G	H	I	J
Particle concentration (µg/ml)	250	500					250	250	250	250
Drug concentration (µg/ml)			5	10	5	10	0.8	1	0.8	1

Fig. S14: MTT assay of ZnFe₂O₄ nanoparticles (A, B), daunorubicin (C, D) and curcumin (E, F), daunorubicin (G-J) and curcumin (K-N) loaded ZnFe₂O₄ nanoparticles treated on normal cell line (CHO).