

Supporting Information

Chitosan hybrid nanoparticles as a theranostic platform for targeted doxorubicin/VEGF shRNA co-delivery and dual-modality fluorescence imaging

Hong Yang^{1,2} §, Min Xu¹ §, Shun Li¹, Xue Shen¹, Tingting Li¹, Jie Yan¹, Chengchen Zhang¹,
Chunhui Wu^{1,2}, Hongjuan Zeng^{1,2}, Yiyao Liu^{1,2*}

¹ Department of Biophysics, School of Life Science and Technology, ² Center for Information in Medicine, University of Electronic Science and Technology of China, Chengdu 610054, Sichuan, P.R. China.

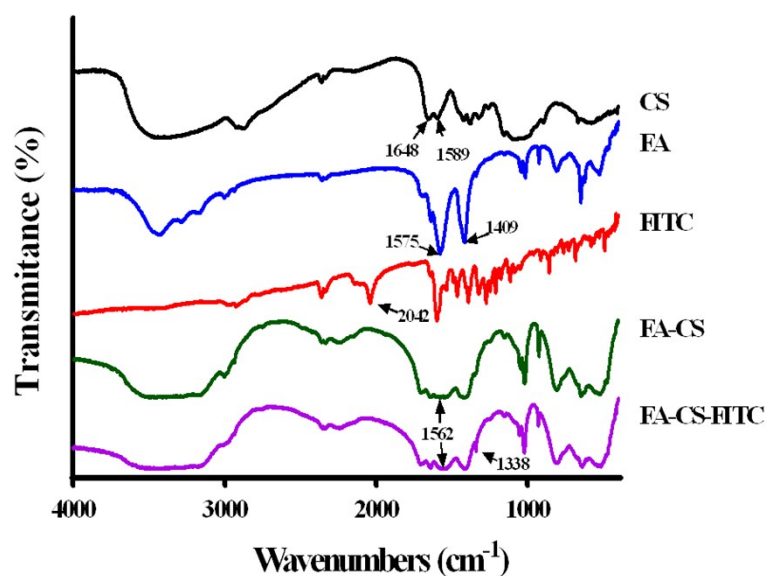


Figure S1. The FTIR spectroscopy of chitosan (CS), folic acid (FA), fluorescein isothiocyanate (FITC), FA-CS, and FA-CS-FITC.

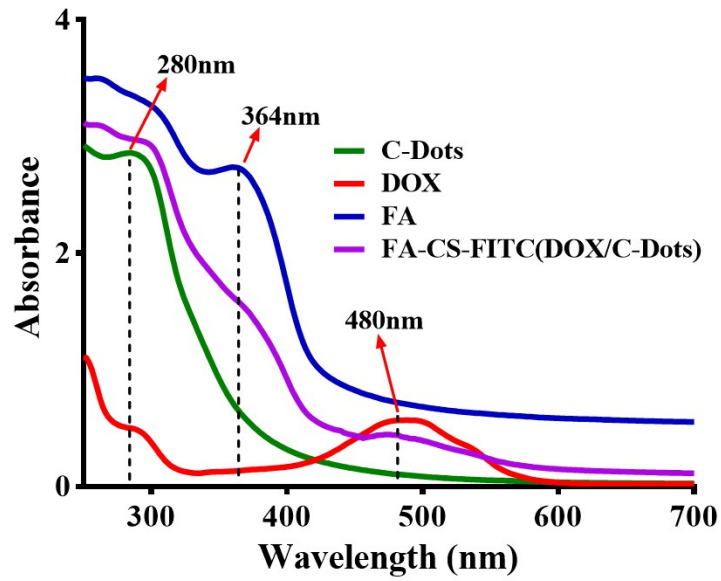


Figure S2. UV-Vis spectra of C-dots, DOX, and FA-CS-FITC(DOX/C-dots).

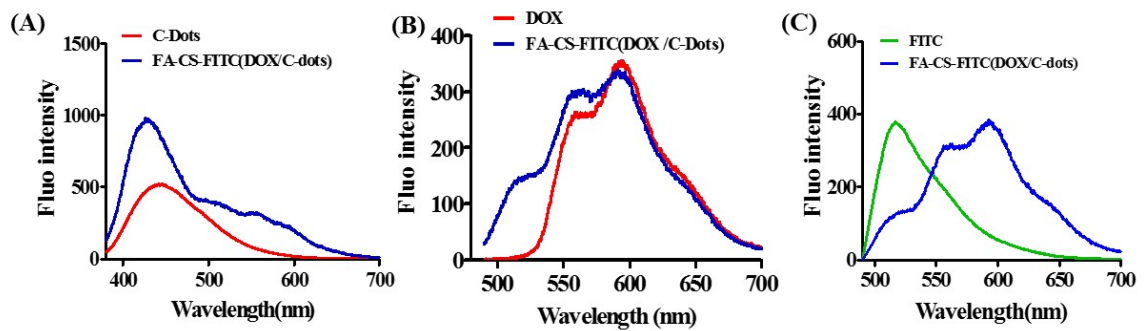


Figure S3. Fluorescence spectrum to detect the presence of C-dots (A), DOX (B) and FITC (C) in the FA-CS-FITC(DOX/C-dots) nanoparticles.

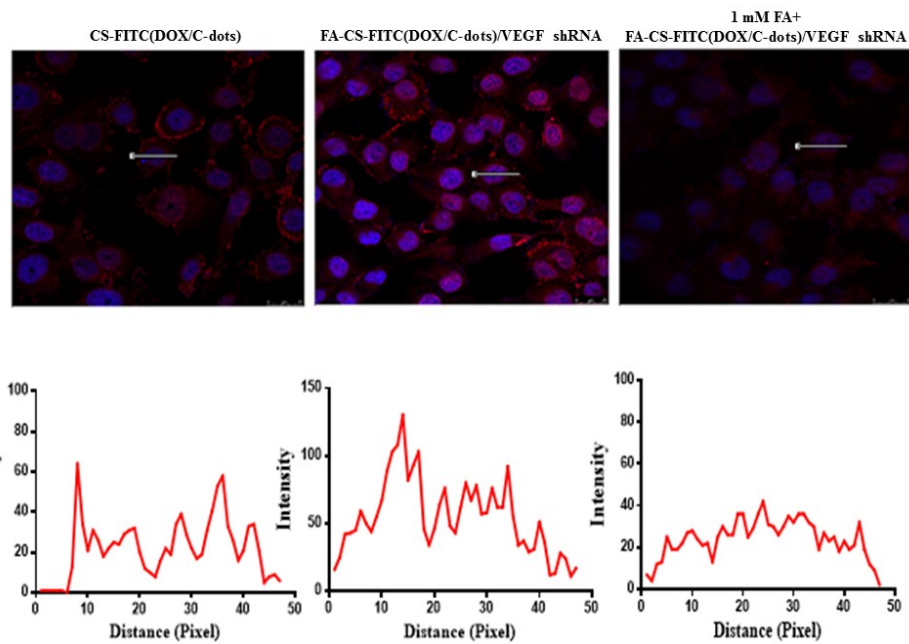


Figure S4. Fluorescence images (up panel) and line-scanning profiles (down panel) of fluorescence intensity for HeLa cells incubated for 6 h with indicated nanoparticles.

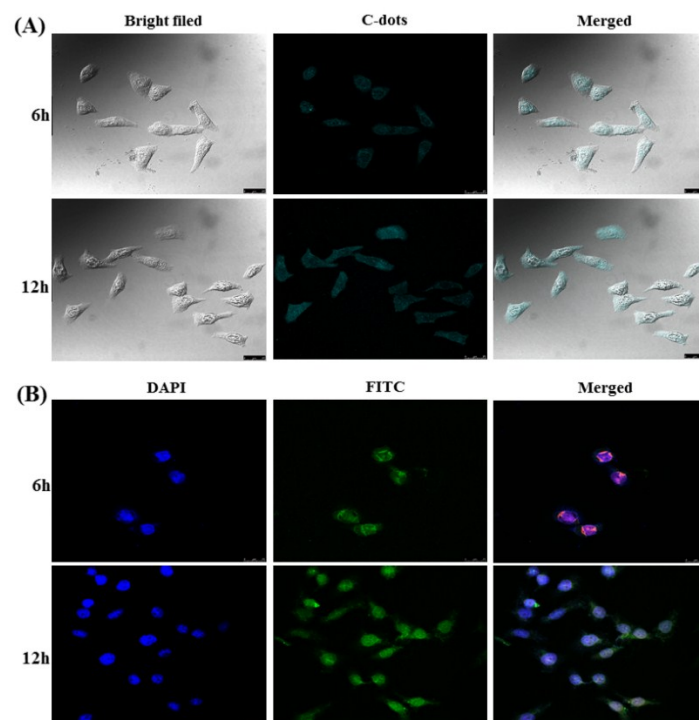


Figure S5. Cellular images of HeLa cells treated with FA-CS-FITC(DOX/C-dots) nanoparticles (20 µg/mL) for 6 h or 12 h by C-dots and FITC detection. Light blue (C-dots), Dark blue (DAPI, nuclei), and green (FITC).