Supporting Information

Carboxymethyl cellulose Grafted Mesoporous Silica hybrid Nanogels for enhanced Cellular uptake and release of Curcumin

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1. SEM of MSNs

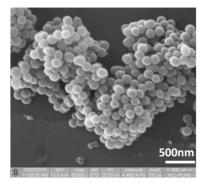


Fig.1S. SEM image of as synthesized MSNs

2. Multimodal distribution of as synthesized and Functionalized MSNs

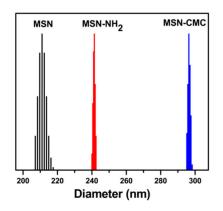


Fig.2S. DLS of MSN, MSN-NH₂ and MSN-CMC

3. Pore diameter of MSN

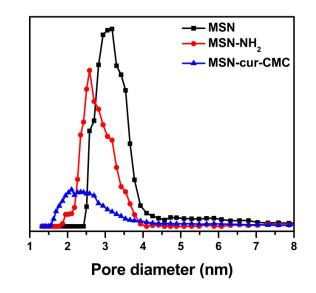
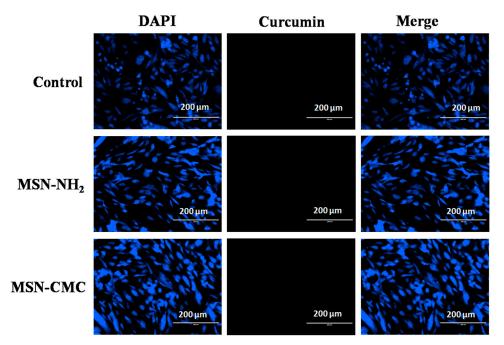


Fig.3S. Pore diameter of MSN, MSN-NH₂ and MSN-cur-CMC using BJH method from N₂-adsortion desorption studies



4. <u>Cellular uptake of MSN-NH₂ and MSN-CMC</u>

Fig.4S. Intracellular uptake of MSN-NH₂ and MSN-CMC using fluorescence microscopy. Images of MDA-MB-231 incubated with 200μ g/ml of MSN-NH₂ and MSN- CMC. Control refers to the non treated MDA-MB-231 cells. Blue fluorescence is due to nuclei staining of cell with DAPI

5. <u>Apoptosis of MSN-NH₂ and MSN-CMC</u>

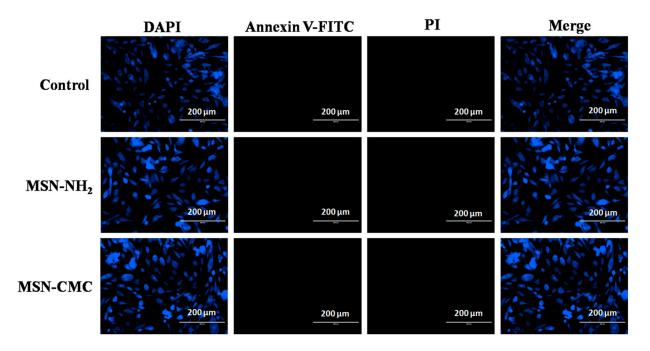
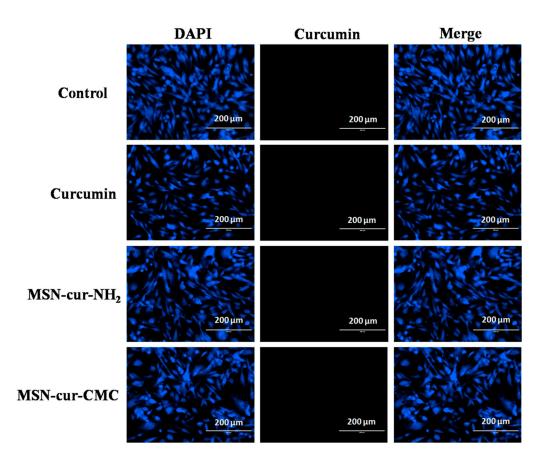


Fig.5S. Apoptosis of MDA-MB-231 cells using fluorescence microscopy. Images are at a magnification of 200 μ m of MDA-MB-231 incubated with 200 μ g/ml of MSN-NH₂ and MSN- CMC. Images of MDA-MB-231 incubated with 200 μ g/ml of MSN-NH₂ and MSN- CMC. Control refers to the non treated MDA-MB-231 cells. Blue fluorescence is due to nuclei staining of cell with DAPI



6. Fluorescence due to curcumin from MSN-cur-NH₂ and MSN-cur-CMC after 48hrs

Fig.6S. Intracellular uptake of $-NH_2$ and -CMC functionalized MSNs using fluorescence microscopy after 48hrs. Images of MDA-MB-231 incubated with 16µg/ml of free curcumin, MSN-cur-NH₂ (GI₅₀=7µg/ml) and MSN-cur-CMC (GI₅₀=1.5µg/ml). Control refers to the non treated MDA-MB-231 cells. Blue fluorescence is due to nuclei staining of cell with DAPI and green due to fluorescence of curcumin release inside the cells.