

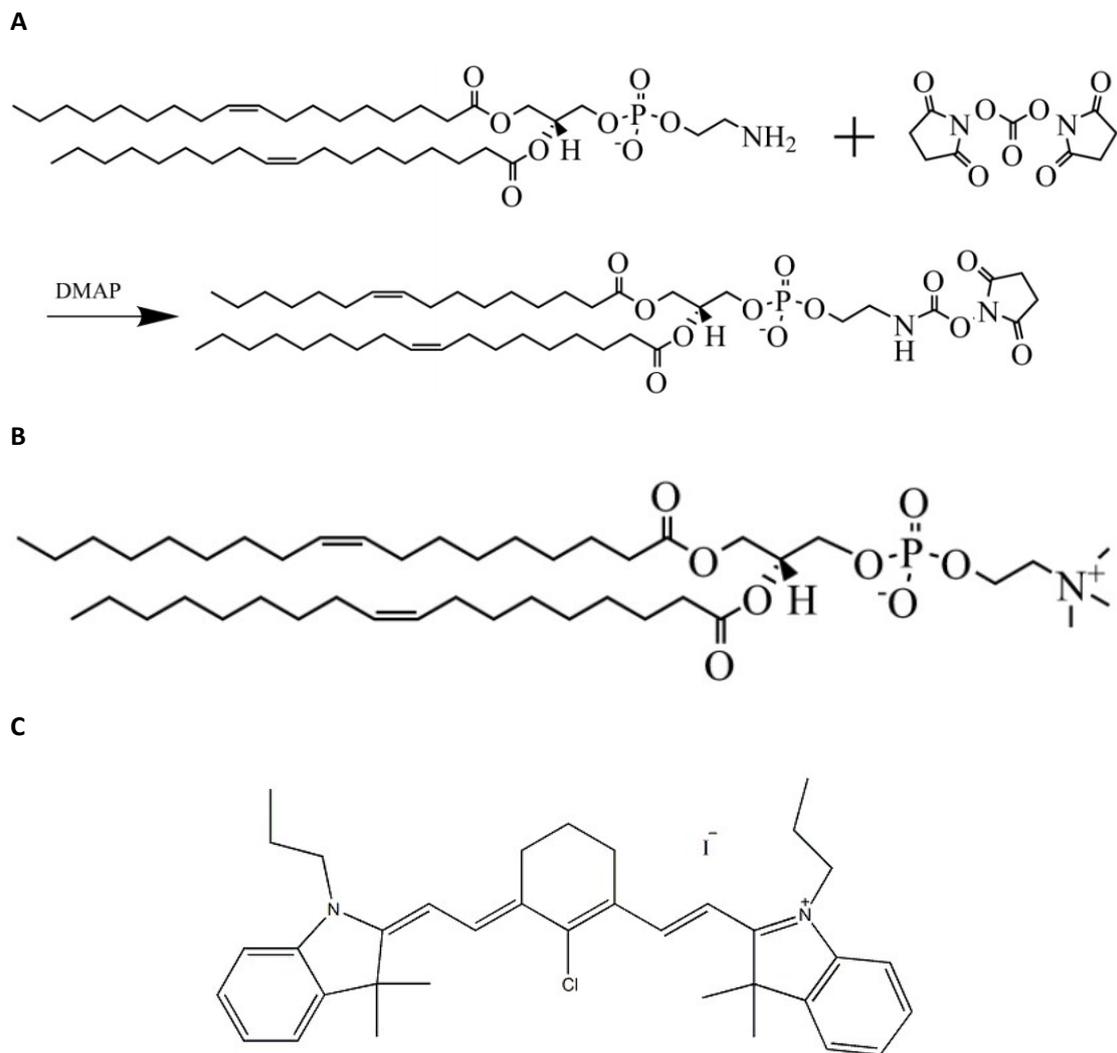
## Supporting Information

### **NIR Light-Activated Dual-Modality Cancer Therapy Mediated by Photochemical Internalization of Porous Nanocarriers with Tethered Lipid Bilayer**

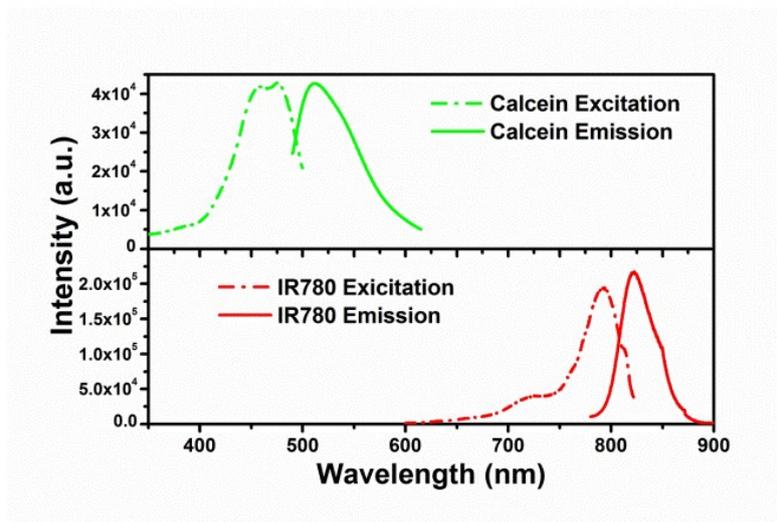
Junjie Liu,<sup>a</sup> Didem Şen Karaman,<sup>b</sup> Jixi Zhang,<sup>\*a</sup> Jessica M. Rosenholm,<sup>b</sup> Xingming Guo<sup>a</sup>  
and Kaiyong Cai<sup>\*a</sup>

a. Key Laboratory of Biorheological Science and Technology, Ministry of Education, College of Bioengineering, Chongqing University, No. 174 Shazheng Road, Chongqing 400044, China. E-mail: jixizhang@cqu.edu.cn, kaiyong\_cai@cqu.edu.cn.

b. Pharmaceutical Sciences Laboratory, Faculty of Science and Engineering, Åbo Akademi University, Tykistökatu 6A, Turku 20520, Finland.

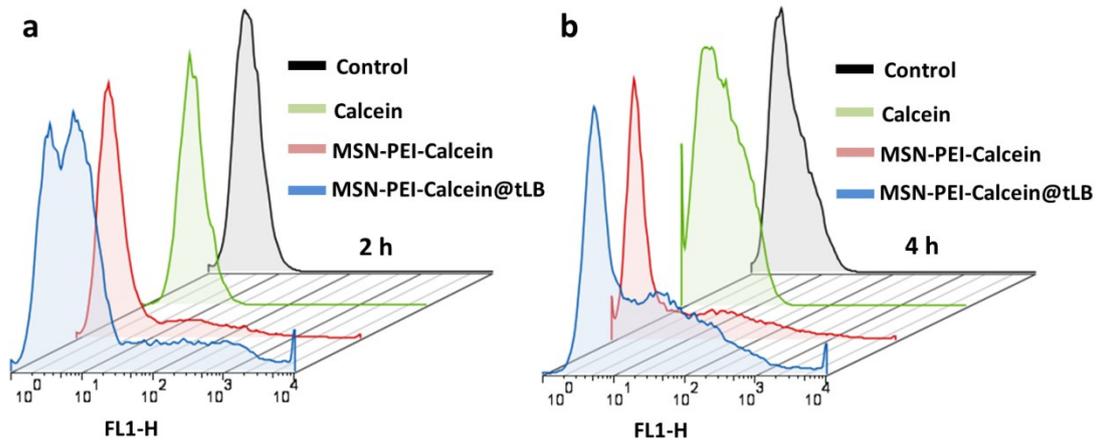


**Scheme S1** (A) Reaction for synthesizing NHS activated DOPE lipid by conjugation between DOPE and coupling agent DSC. Molecular structures of DOPC (B) and IR-780 iodide (C).

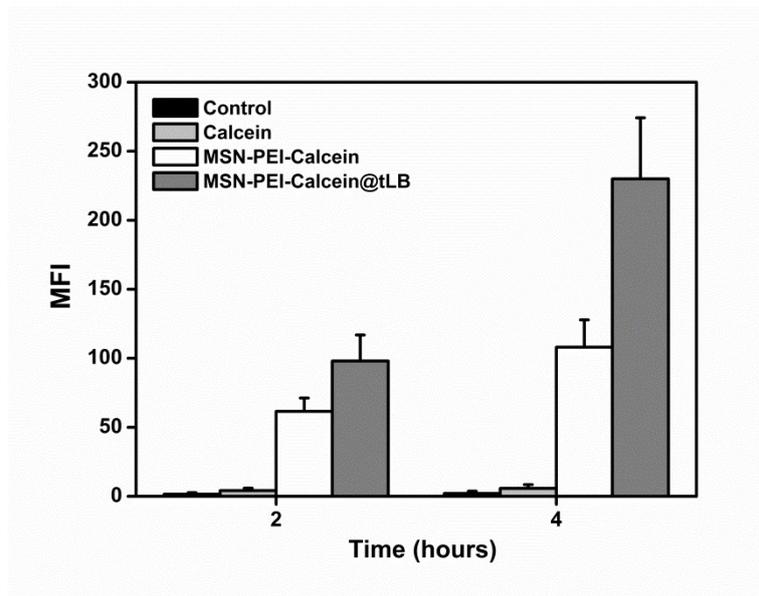


**Fig. S1** Excitation and emission spectra of MSN-PEI-calcein@tLB-IR780.

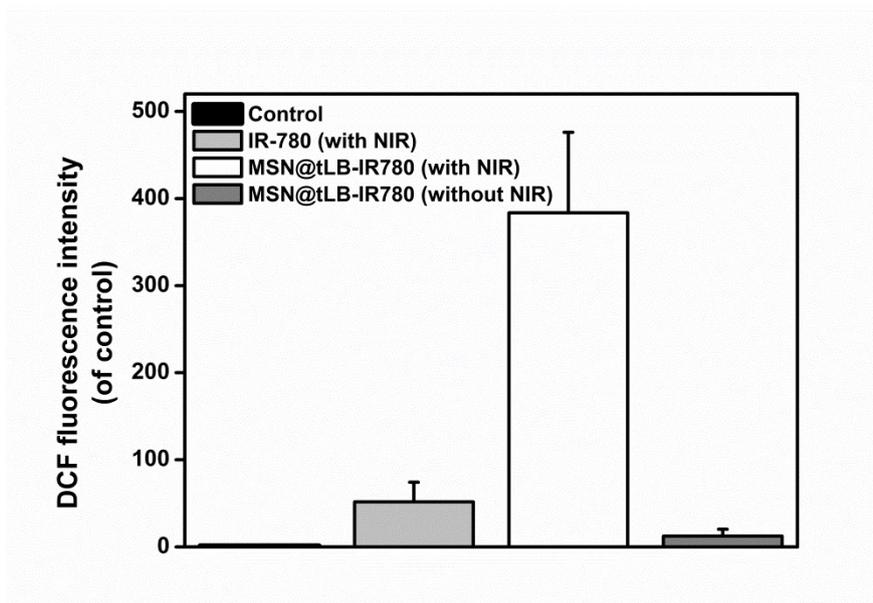
**A**



**B**



**Fig. S2 (A)** Representative FACS analysis image show the calcein-positive percentage of MCF-7 cells treated with free calcein, MSN-PEI-calcein and MSN-PEI-calcein@tLB for 2 and 4 h, respectively, where MCF-7 cells without treatment act as control. **(B)** Quantitative mean fluorescence intensity (MFI) analysis after cells with different treatments for 2 and 4 h.



**Fig. S3** Detection of ROS production. Relative DCF fluorescence intensity after cells treated with free IR-780 (with irradiation), MSN@tLB-IR780 (with or without irradiation) compare to control group. The control group was performed that cells without any treatment. Irradiation condition: the cells were exposed to 808 nm laser ( $1.2 \text{ W/cm}^2$ ) for 10 min.

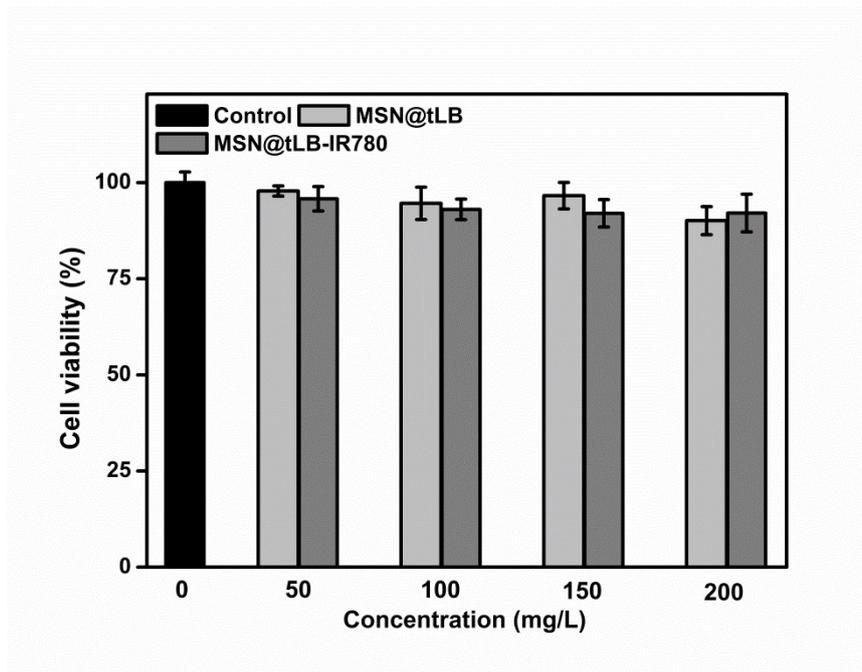


Fig. S4 Cytotoxicity of formulations with different concentrations on MCF-7 cells in dark.