## **Supporting Information**

## Sorafenib and tetrakis (4-carboxyphenyl) porphyrin assembled nanoparticles for synergistic targeted chemotherapy and sonodynamic therapy of hepatocellular carcinoma

Yongzhi Chen,<sup>a</sup> Qiuxia Tan,<sup>b</sup> Yuanyu Tang,<sup>b</sup> E Pang,<sup>b</sup> Rui Peng,<sup>a</sup> Minhuan Lan,<sup>\*b</sup> Dousheng Bai<sup>\*a</sup>

<sup>*a*</sup> Department of Hepatobiliary Surgery, Clinical Medical College, Yangzhou University, 98 West Nantong Rd, Yangzhou 225000, P.R. China.

<sup>b</sup> Hunan Provincial Key Laboratory of Micro & Nano Materials Interface Science, College of Chemistry and Chemical Engineering, Central South University, Changsha, Hunan, 410083, P. R. China.

Corresponding authors.

minhuanlan@csu.edu.cn (M. Lan); drbaidousheng@yzu.edu.cn (D. Bai)



**Fig. S1** The absorbance and concentration relationship of TCPP. (Abs represents absorbance of STP NPs, C represents the concentration of STP NPs)



Fig. S2 High Performance Liquid Chromatography (HPLC) of STP NPs.



Fig. S3 SEM images of STP NPs at days 3 (A), 5 (B), and 7 (C).



Fig. S4 Time-dependent particle size variation and SEM of SP NPs and TP NPs.



Fig. S5 UV-vis absorption spectra of STP NPs after US irradiation for different times.



Fig. S6 UV-vis absorption spectra of ABDA-Na2 in the absence of STP NPs after US

irradiation for different times.



Fig. S7 Inverted microscope images displaying intracellular uptake of STP NPs (red



**Fig. S8** Incubation time-dependent fluorescent images of HepG2 cells in the presence of SOR-TCPP@PEG NPs.

fluorescence).



Fig. S9 Fluorescent images of O22-HepG2 cells under different treatments: SP, SP +

US, TP, and TP + US.



Fig. S10 Calcein-AM- and PI-stained HepG2 cells after different treatments: SP, SP +

US, TP, and TP + US.



Fig. S11 The full gel and blot image of western blot.



Fig. S12 (A) Fluorescence imaging of SOR-TCPP@PEG NPs within mouse tumor at

8 h. (B) Tumor photographs of nude mice were taken after TP + US treatments.



**Fig. S13** Western blotting was used to analyze the expression of VEGFR-2 in tumor tissues after different treatments.