

SUPPORTING INFORMATION

Bifunctional pH-Sensitive Zn(II)-Curcumin Nanoparticles/siRNA Effectively Inhibit Growth of Human Bladder Cancer Cells *in Vitro* and *in Vivo*

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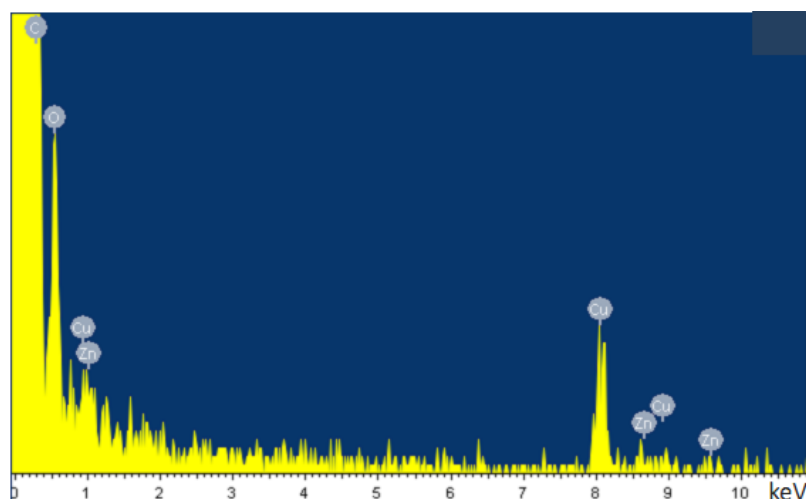


Fig. S1 EDS spectrum of the obtained Zn(II)-Cur NPs. Cu peak came from the TEM grid.

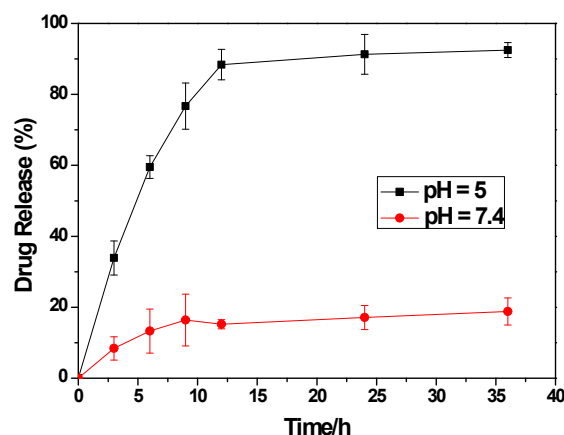


Fig. S2 Drug release profiles under pH control at two different pH = 5 and 7.4 (n = 3, *p < 0.01).

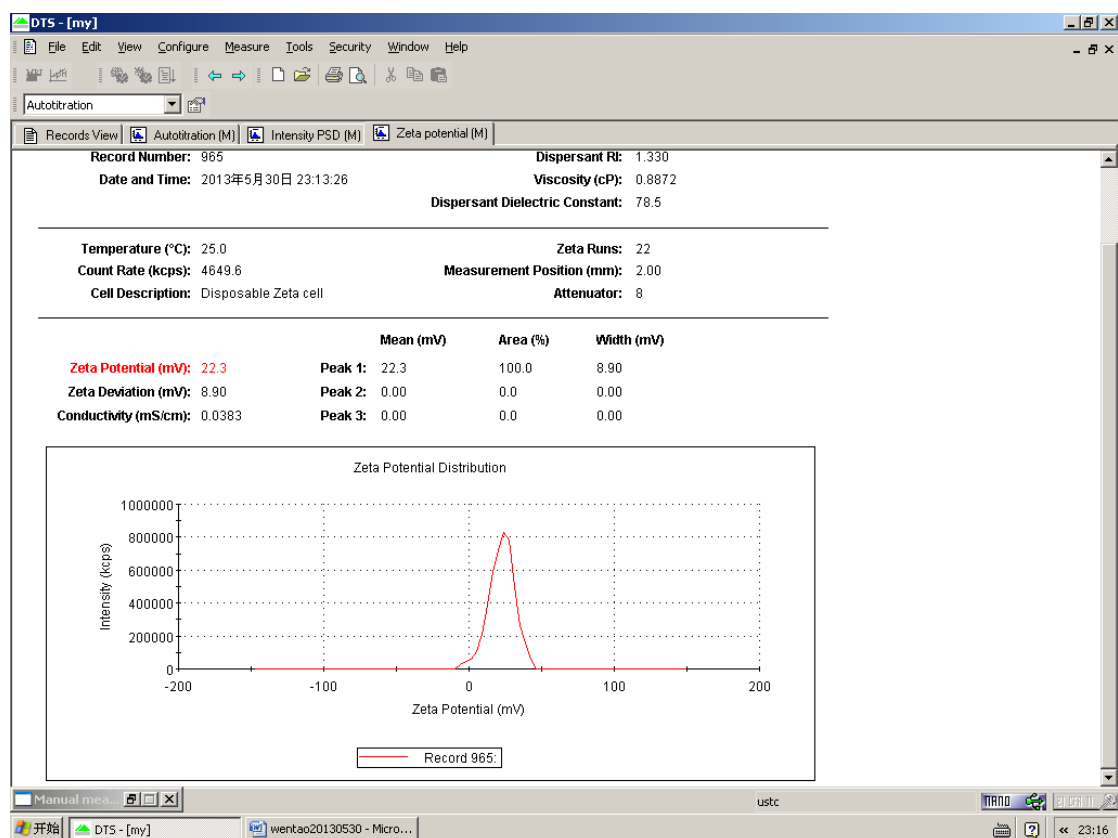


Fig. S3 Zeta potential distribution of Zn(II)-Cur NPs.

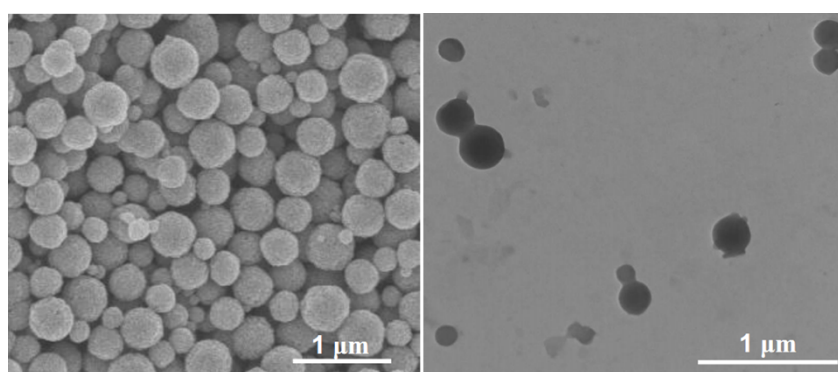


Fig. S4 SEM (left) and TEM (right) image of Zn(II)-Cur NPs/siEIF5A2 complex.

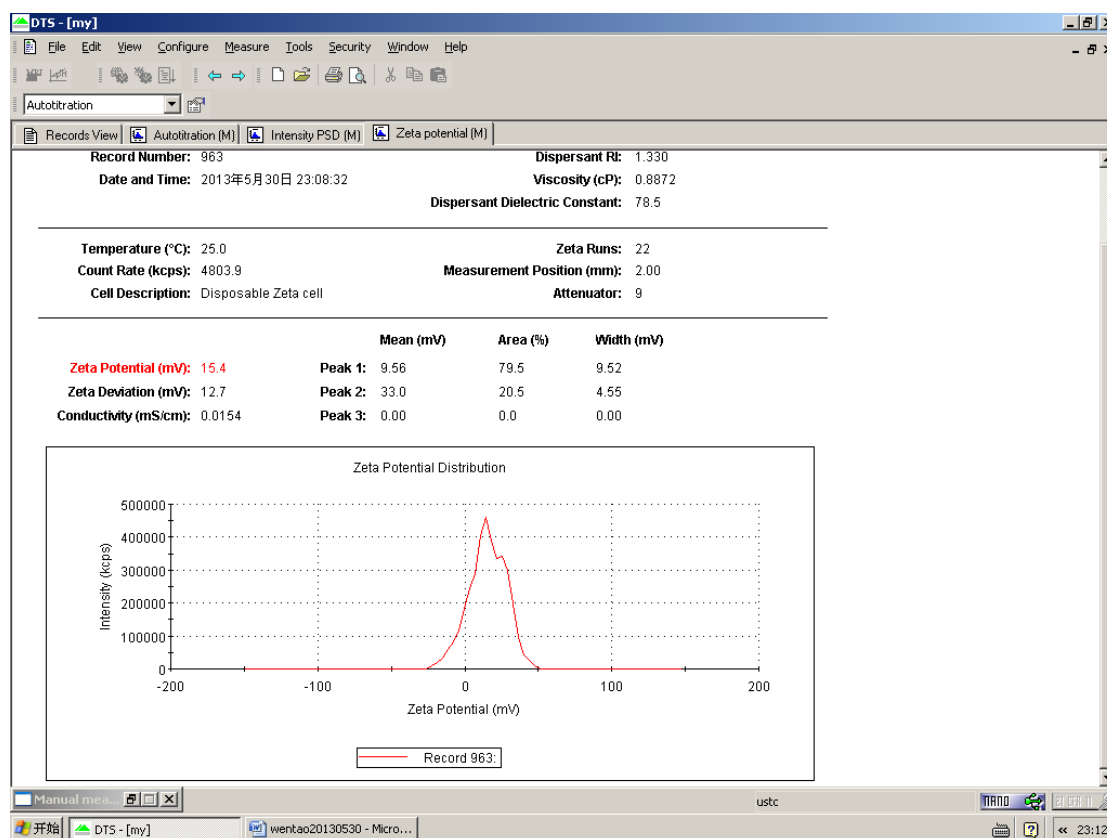


Fig. S5 Zeta potential distribution of Zn(II)-Cur NPs/siEIF5A2 complex.

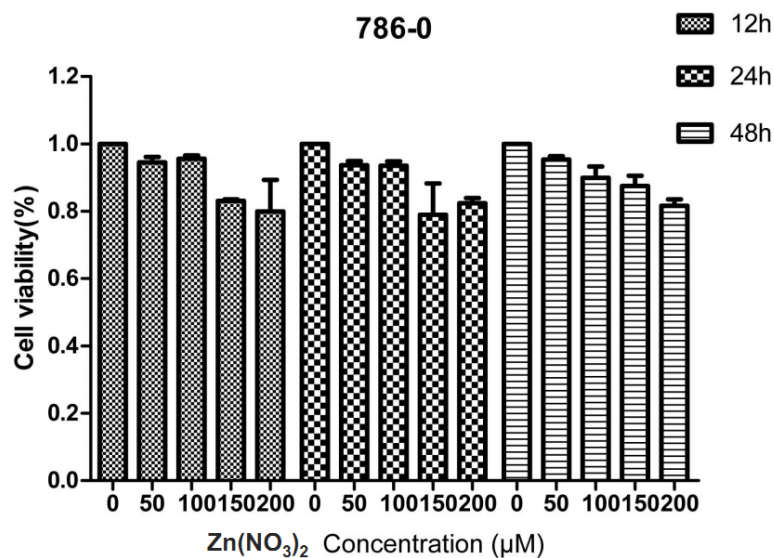


Fig. S6 The toxicity of Zn²⁺ at the concentration of 0, 50 nM, 100 nM, 150 nM and 200 nM at 12 h, 24 h and 48 h (n = 6, *P < 0.05, **P < 0.01, and ***P < 0.001 compared to other groups).

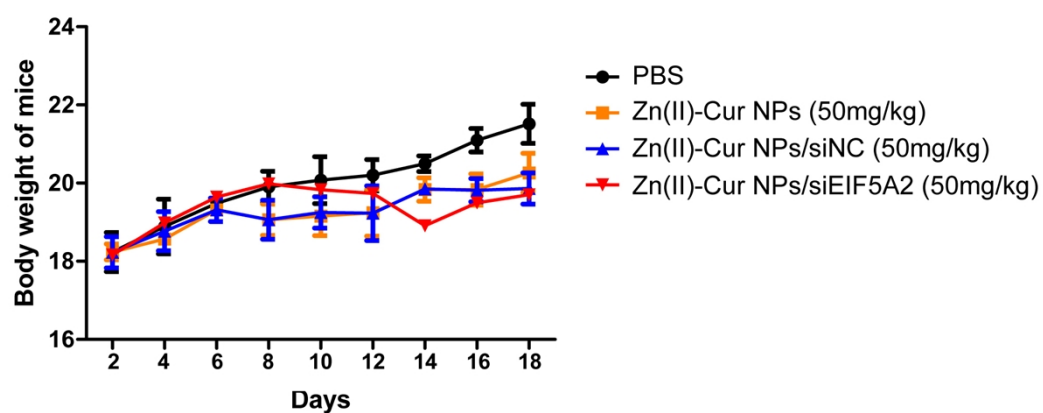


Fig. S7 Changes of body weight after injection of PBS, Zn(II)-Cur NPs, Zn(II)-Cur NPs/siNC and Zn(II)-Cur NPs/siEIF5A2 at 50 mg/kg. The means $\pm$ SD are cumulative results from five independent experiments.