Supporting Information

Formation of Platinum Nanocrystals on Silicon

Nanotubes and Corresponding Anti-Cancer Activity *In*

Vitro

Nguyen T. Le, ¹ Giridhar R. Akkaraju, ² Jeffery L. Coffer^{1*}

¹ Department of Chemistry and Biochemistry, Texas Christian University, Fort Worth, TX, 76129,

USA

e-mail: j.coffer@tcu.edu

² Department of Biology, Texas Christian University, Fort Worth, TX, 76129, USA

Figure S1. Structural morphology of pSiNTs after functionalization with APTES.

Figure S2. Shell thickness of Unmodified pSiNTs and APTES-pSiNTs.

Figure S3. Pt NCs-pSiNTs after incubation in different K₂PtCl₄ salt concentrations and time

Figure S4. Pt NCs formed on unmodified pSiNTs after 24 h.

Figure S5. Pt NCs formed on unmodified SiO₂ NTs and APTES-functionalized SiO₂ NTs.

Figure S6. Normalized toxicity based on wt% Pt

Figure S7. Citrate-capped Pt NCs.

Figure S8. Cytotoxicity of Pt NCs $(3.5 \pm 1.1 \text{ nm})$ and K_2 PtCl₄ at different doses.

Figure S9. Localization of Pt NCs-pSiNTs in HeLa cells

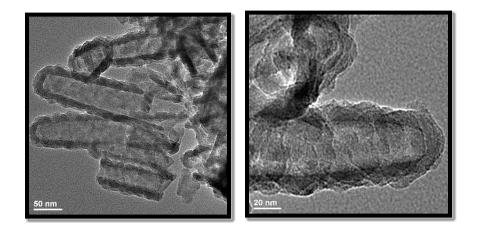


Figure S1. Structural morphology of pSiNTs after functionalization with APTES.

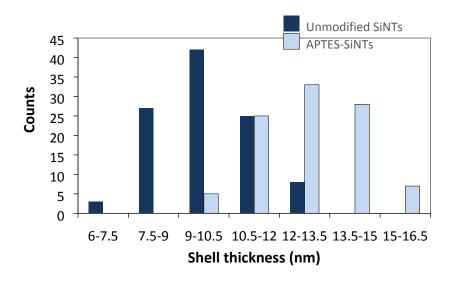


Figure S2. Shell thickness of Unmodified pSiNTs and APTES-pSiNTs.

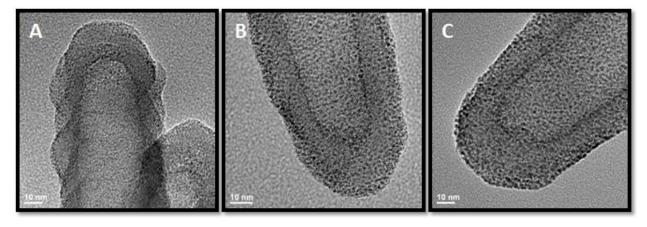


Figure S3. Pt NCs-pSiNTs after incubation in different K_2PtCl_4 salt concentrations and time A) $K_2PtCl_4 = 0.5 \text{ mM } (4 \text{ h}), B) K_2PtCl_4 = 0.5 \text{ mM } (24 \text{ h}), C) K_2PtCl_4 = 3.3 \text{ mM } (24 \text{ h}). Scale bar = 10 \text{ nm}.$

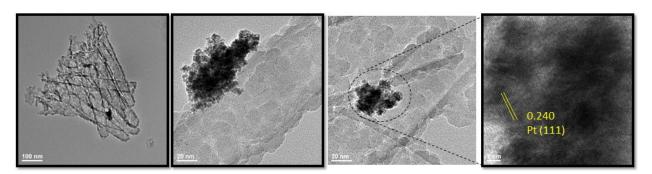


Figure S4. Pt NCs formed on unmodified pSiNTs after 24 h.

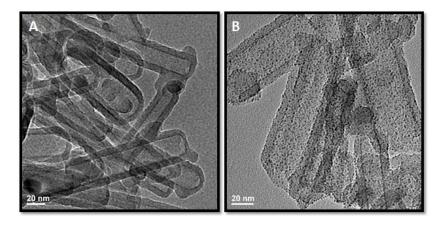


Figure S5. Pt NCs formed on A) unmodified SiO_2 NTs and B) APTES-functionalized SiO_2 NTs.

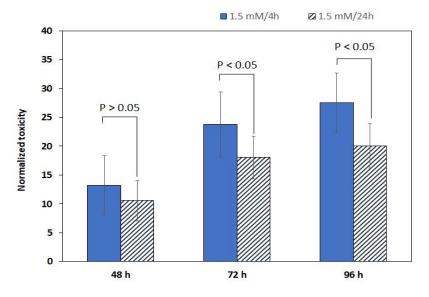


Figure S6. Normalized toxicity based on wt% Pt.

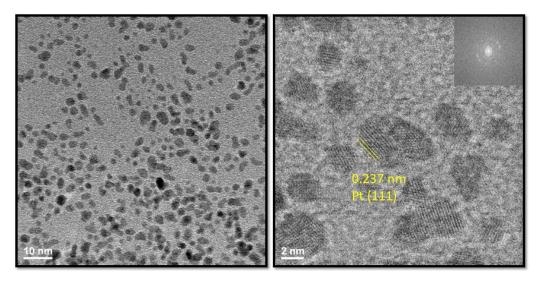


Figure S7. Citrate-capped Pt NCs.

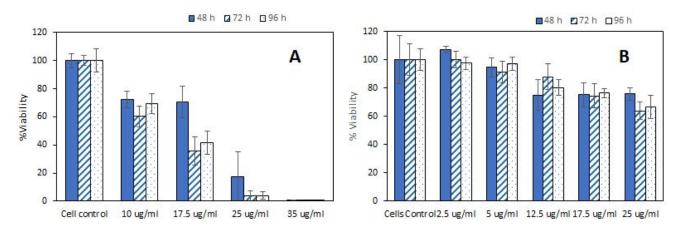


Figure S8. Cytotoxicity of A) Pt NCs $(3.5 \pm 1.1 \text{ nm})$ and B) K_2PtCl_4 at different doses.

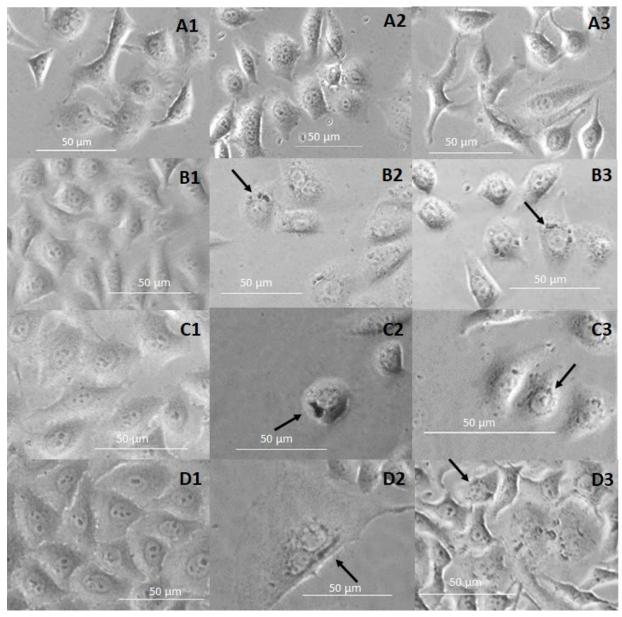


Figure S9. Localization of Pt NCs-pSiNTs in HeLa cells after 4h (A1-A3), 24h (B1-B3), 48h (C1-C3) and 72h (D1-D3) under brightfield of fluorescence microscope (non-fluorescence method). A1-D1: Cell with no treatment; A2-D2: 1.5 mM/4h (30-35 %wt Pt) Pt NCs-pSiNTs; A3-D3: 1.5 mM/24h (40-45 %wt Pt) Pt NCs-pSiNTs.