Supplementary Material for

Three Pt(II) complexes based on Thiosemicarbazone: synthesis, HSA interaction, cytotoxicity, apoptosis and cell cycle arrest

Xu-Dong Lin,^{1a} Ya-Hong Liu,^{1a} Cheng-Zhi Xie,^a Wei-Guo Bao,^a Jun Shen^{b,*} and Jing-Yuan Xu^{a,*}

^a Department of Chemical Biology and Tianjin Key Laboratory on Technologies Enabling Development of Clinical Therapeutics and Diagnostics (Theranostics), School of Pharmacy, Tianjin Medical University, Tianjin 300070, P. R. China.

^bDepartment of Sanitary Chemistry, School of Public Health, Tianjin Medical University, Tianjin 300070, P. R. China.

*Corresponding author:

Tel: +86 022 83336658, E-mail: xujingyuan@tmu.edu.cn (Prof. J. Y. Xu); Tel: +86 022 83336638, E-mail: shenjun@tmu.edu.cn (Prof. J. Shen).



Fig. S1-1. ¹H NMR spectrum of Ligand 1.



Fig. S1-2. ¹³C NMR spectrum of Ligand 1.



Fig. S1-3. ¹H NMR spectrum of Ligand 2.



Fig. S1-4. ¹³C NMR spectrum of Ligand 2.



Fig. S1-5. ¹H NMR spectrum of Ligand 3.



Fig. S1-6. ¹³C NMR spectrum of Ligand 3.



Fig. S2-1. Fluorescence spectra of HSA in the different concentrations of **1** at room temperature.



Fig. S2-2. Fluorescence spectra of HSA in the different concentrations of **2** at room temperature.



Fig. S2-3. UV-vis absorption spectra of HSA in the absence and presence of 1.



Fig. S2-4. UV-vis absorption spectra of HSA in the absence and presence of 2.



Fig. S2-5. Synchronous fluorescence spectra of HSA upon addition of 1.



Fig. S2-6. Synchronous fluorescence spectra of HSA upon addition of 2.



Fig. S3-1. Hoechst 33342 staining detected apoptosis in HeLa cells after treatment by 1 for 48 h at the concentrations of 0, 2.5, 5, and 10 μ M.



Fig. S3-2. Hoechst 33342 staining detected apoptosis in HeLa cells after treatment by 2 for 48 h at the concentrations of 0, 2.5, 5, and 10 μ M.



Fig. S4-1. The quantitative analysis of the proportion of viable, necrosis and apoptosis cells against HeLa cells after the treatment with 1 for 48 h at concentration of 0, 2.5, 5, and 10 μ M.



Fig. S4-2. The quantitative analysis of the proportion of viable, necrosis and apoptosis cells against HeLa cells after the treatment with 2 for 48 h at concentration of 0, 2.5, 5, and 10 μ M.



Fig. S5. Clonogenic assay (crystal violet staining) in HeLa cells exposed to varying concentrations of **3**.