

## **Supporting Information**

### **Investigation on New Ruthenium(II) Hydrazone Complexes as Anticancer Agent and Their Interaction with Biomolecules**

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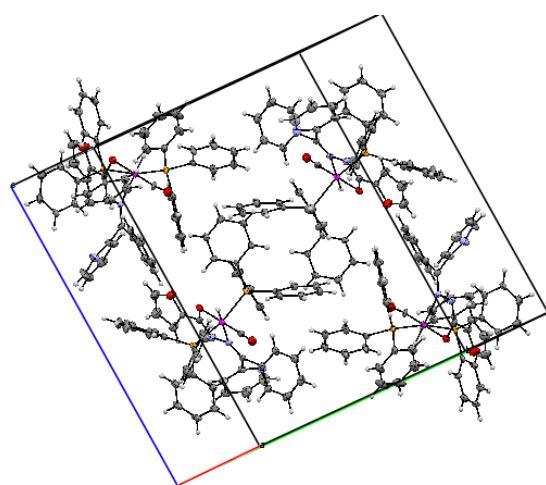
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Electronic supplementary information (ESI) available: CCDC reference number of the complex is 917920, 917921. For crystallographic data in CIF see DOI:

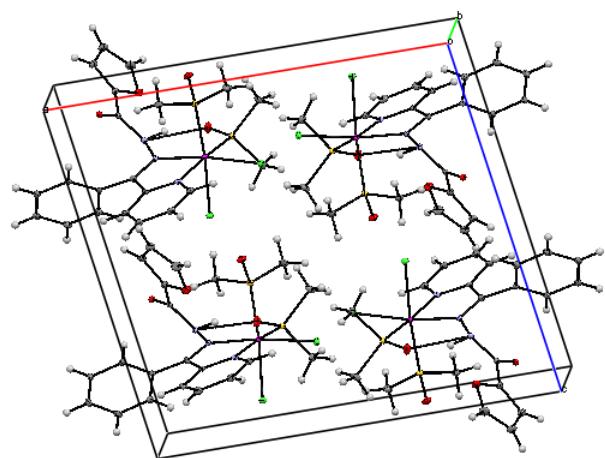
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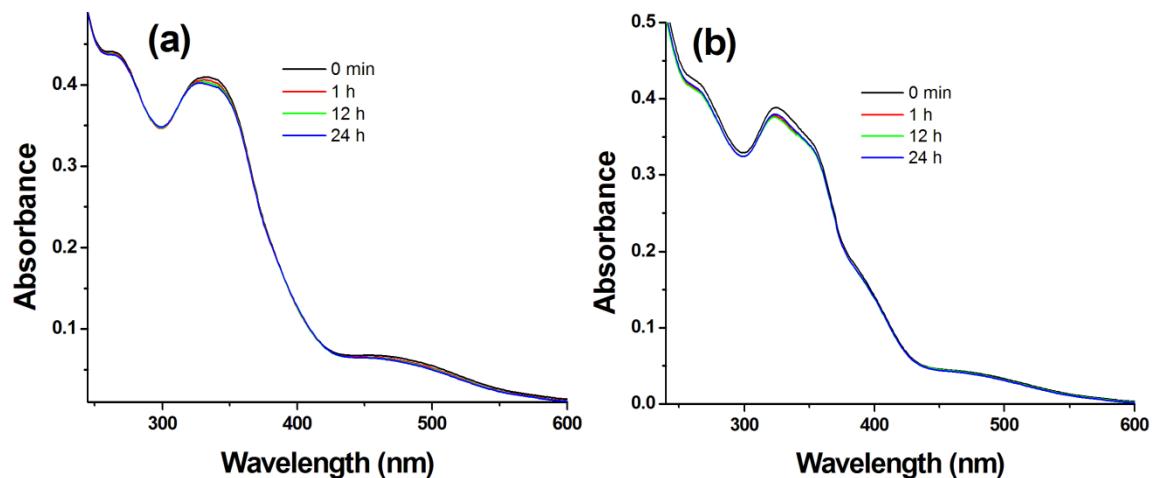


**Complex 1**



**Complex 2**

**Figure S1** Packing diagram of Ru(II) complex with inter- molecular hydrogen bonding



**Figure S2** UV-Vis. absorption spectrum of complexes **1** and **2** in aqueous PBS buffer (phosphate buffered saline solution), at pH 7.4, to a final concentration of  $1 \times 10^{-3}$  M recorded after different time intervals.

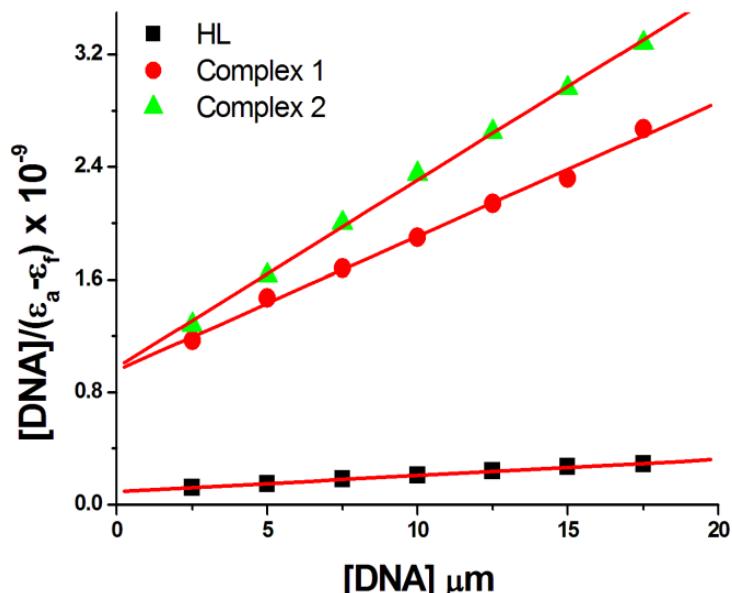


Figure S3 Plots of  $[DNA]/(\epsilon_a - \epsilon_f)$  versus  $[DNA]$  for the compounds with CT-DNA.

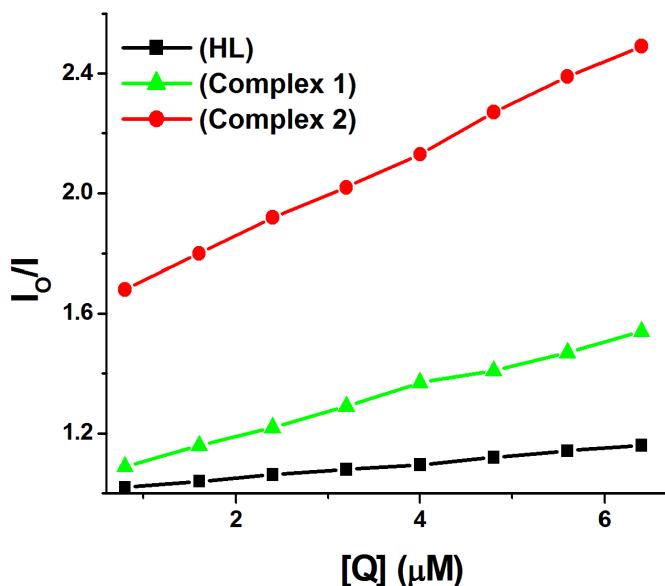


Figure S4 Stern–Volmer plots of the fluorescence titrations of the ligand and the complexes.

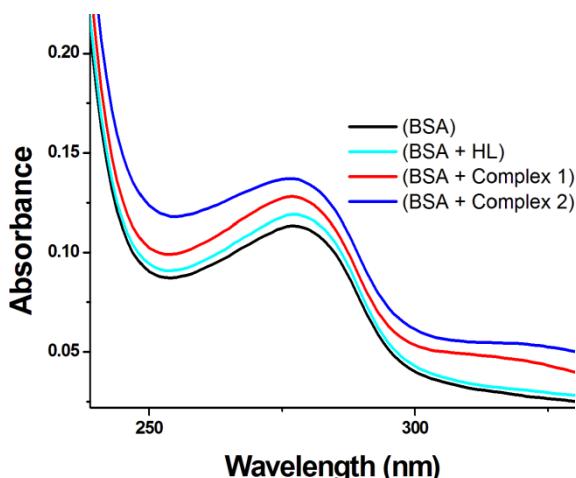
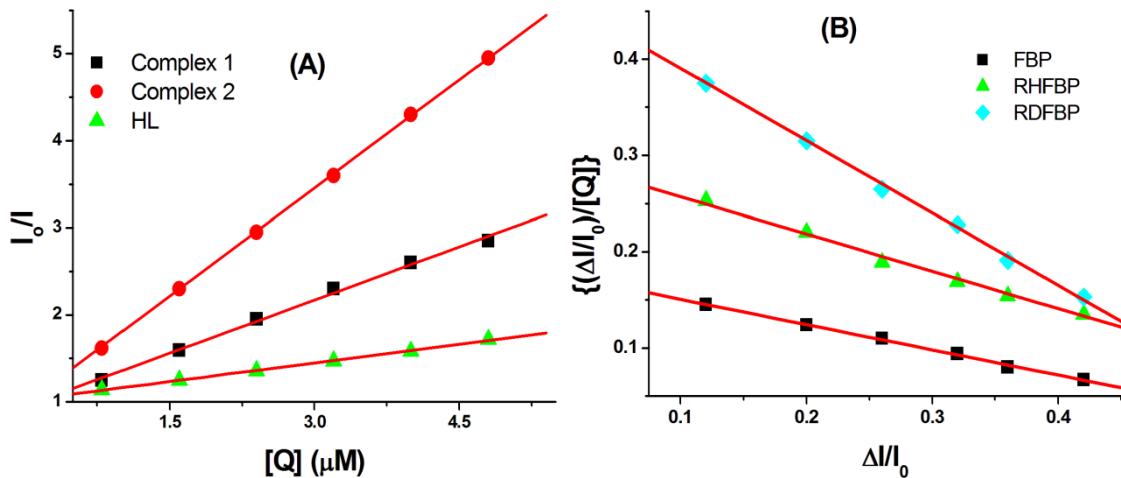
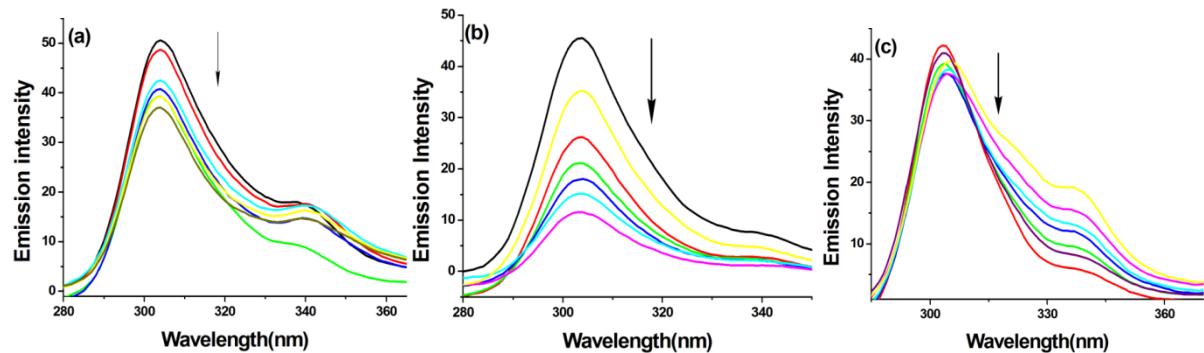


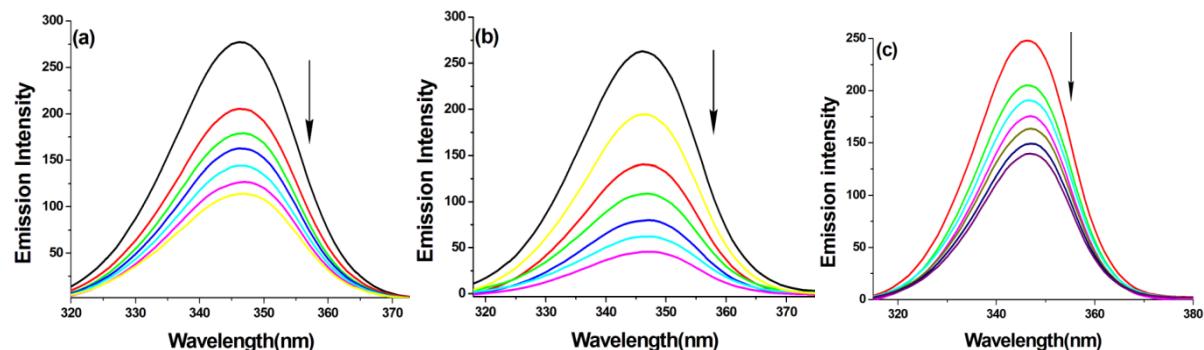
Figure S5 Electronic absorption spectra of BSA (10  $\mu M$ ), with ligand and complexes



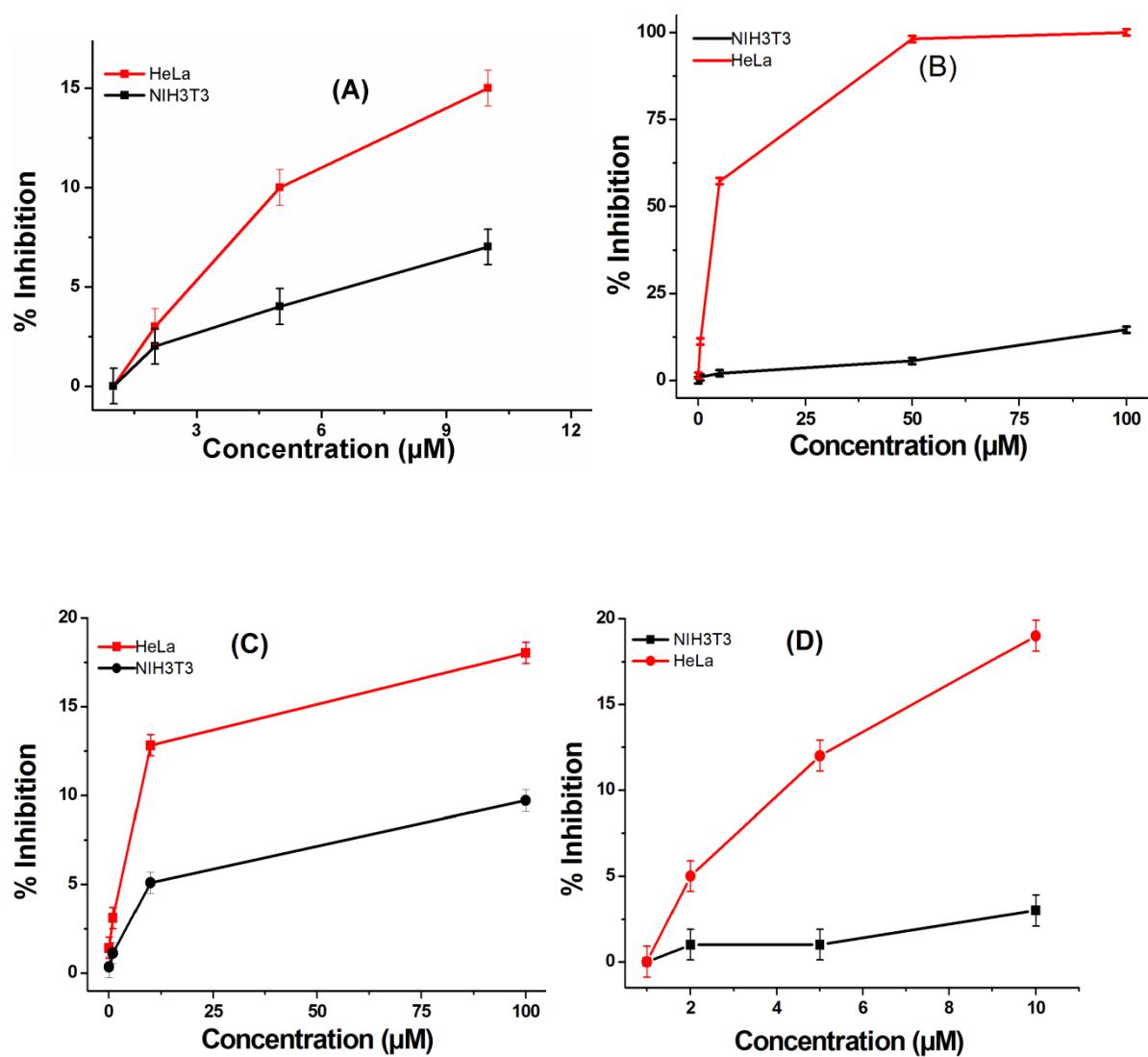
**Figure S6** Stern–Volmer plots (A) and Scatchard plots (B) of the fluorescence titration of the ligand and the complexes with BSA.



**Figure S7** Synchronous spectra of BSA (1  $\mu\text{M}$ ) in the presence of increasing amounts of the complexes **1** (a) and **2**(b) and ligand **HL**(c) (0-12  $\mu\text{M}$ ) at a wavelength difference of  $\Delta\lambda = 15$  nm. The arrow shows the emission intensity decreases upon the increase in concentration of the compounds.



**Figure S8** Synchronous spectra of BSA (1  $\mu\text{M}$ ) in the presence of increasing amounts of the complexes **1** (a) and **2**(b) and the ligand **HL**(c) (0-12  $\mu\text{M}$ ) at a wavelength difference of  $\Delta\lambda = 60$  nm. The arrow shows the emission intensity decreases upon the increase in concentration of the compounds.



**Figure S9** Cytotoxicity of complex 1**(A)**, complex 2 **(B)**, Ligand **(C)** and *cisplatin* **(D)**, respectively