

Electronic Supplementary Material (ESI) for RSC Advances

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Supplementary Materials

Photophysical, G-quadruplex DNA binding and cytotoxic properties of terpyridine complexes with naphthalimide ligand

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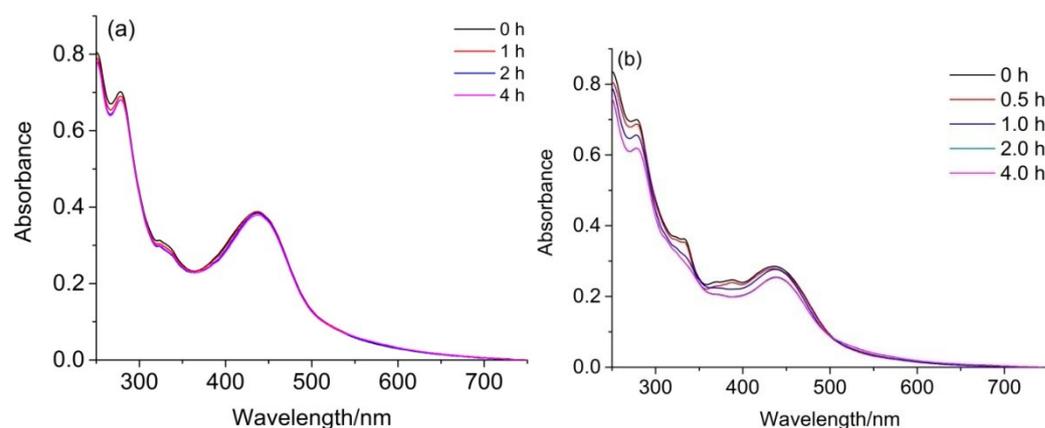


Fig. S1 UV-Vis absorption spectral changes of **3** (30 μ M) in HEPES buffer (10 mM, pH 7.4) incubated in the presence of (a) 0.15 M KCl and (b) 1 mM glutathione.

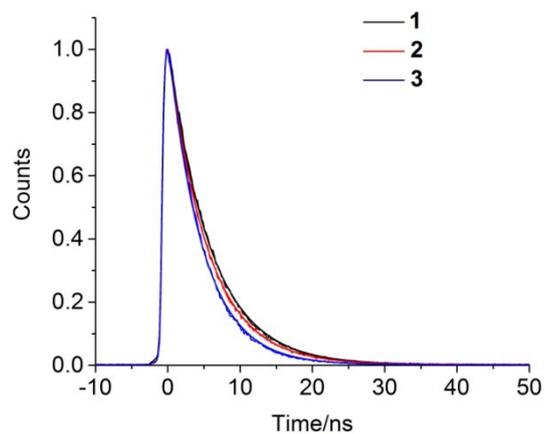


Fig. S2 Fluorescence decay profiles of **1-3** (4 μM) in HEPES buffer solution (10 mM, pH 7.4) containing 0.1 M KCl ($\lambda_{\text{ex}} = 435 \text{ nm}$).

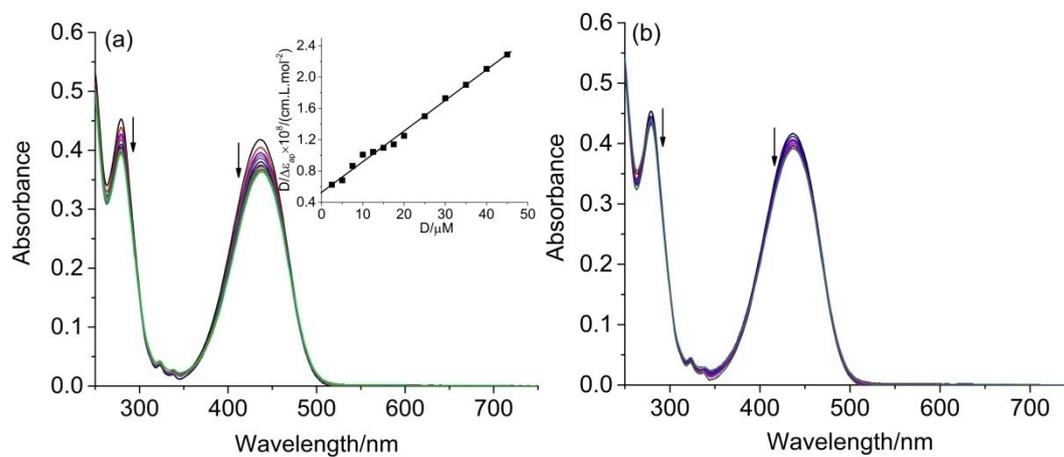


Fig. S3 UV-Vis titration of **1** (30 μM) with (a) CT DNA and (b) Htelo in HEPES (10 mM, pH 7.4) containing 0.1 M NaCl. The arrow indicates the changes upon addition of DNA. Inset: Plot of $D/\Delta\epsilon_{\text{ap}}$ versus D .

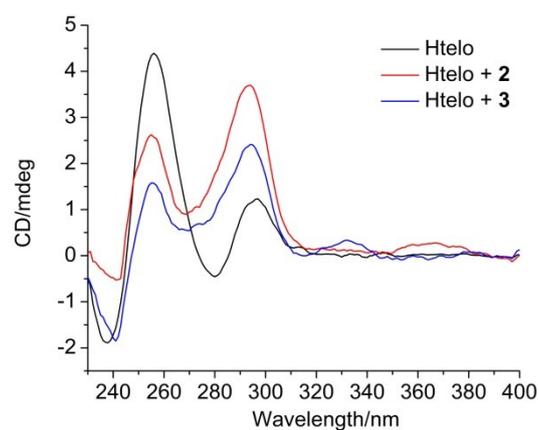


Fig. S4 CD spectra of Htelo (10 μM) in 10 mM Tris-HCl (pH 7.4) in the absence of monovalent cation upon addition of the complexes **2-3** (20 μM).

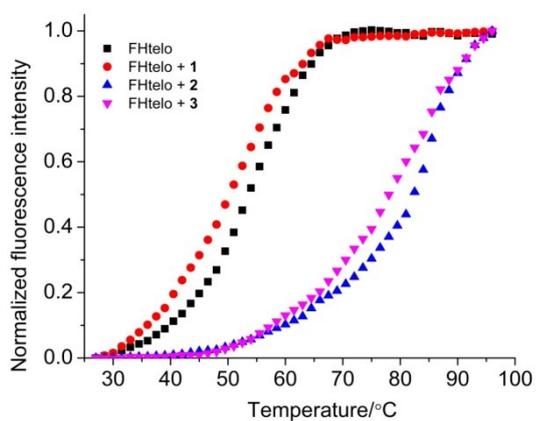


Fig. S5 FRET melting assay of the labeled FHtelo (0.2 μM) in lithium cacodylate buffer (10 mM, pH 7.2), containing KCl (10 mM) and LiCl (90 mM), in the presence of **1-3** (1.0 μM).

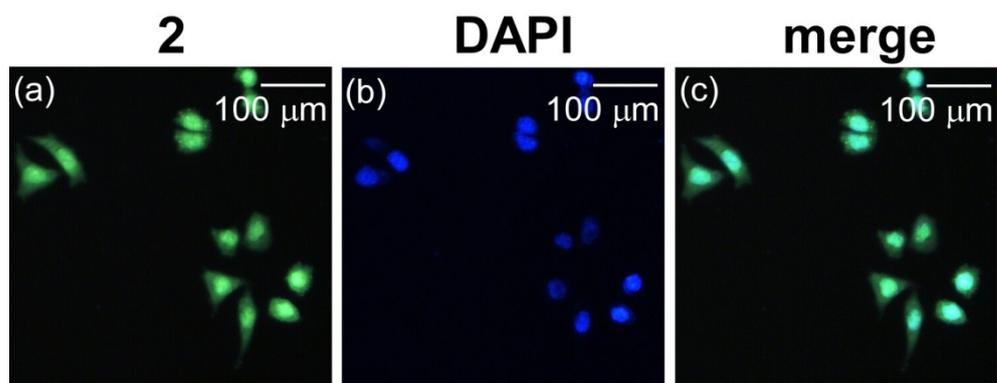


Fig. S6 Double staining of A549 cells with **2** (5 μM) and DAPI (5 μM) at 37 $^{\circ}\text{C}$ for 0.5 h (a-c). Images from left to right: fluorescence image of **2** (laser 488 nm); laser images of DAPI (laser 405 nm); merged image.

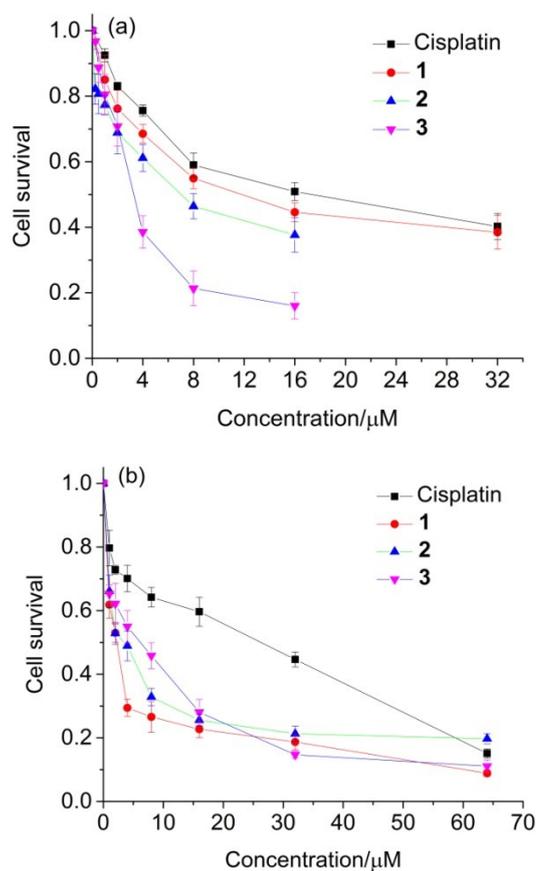


Fig. S7 The cytotoxic activities of **1**, **2**, **3** and cisplatin against (a) A549 and (b) NIH3T3 cells. The drug treatment period is 48 h.

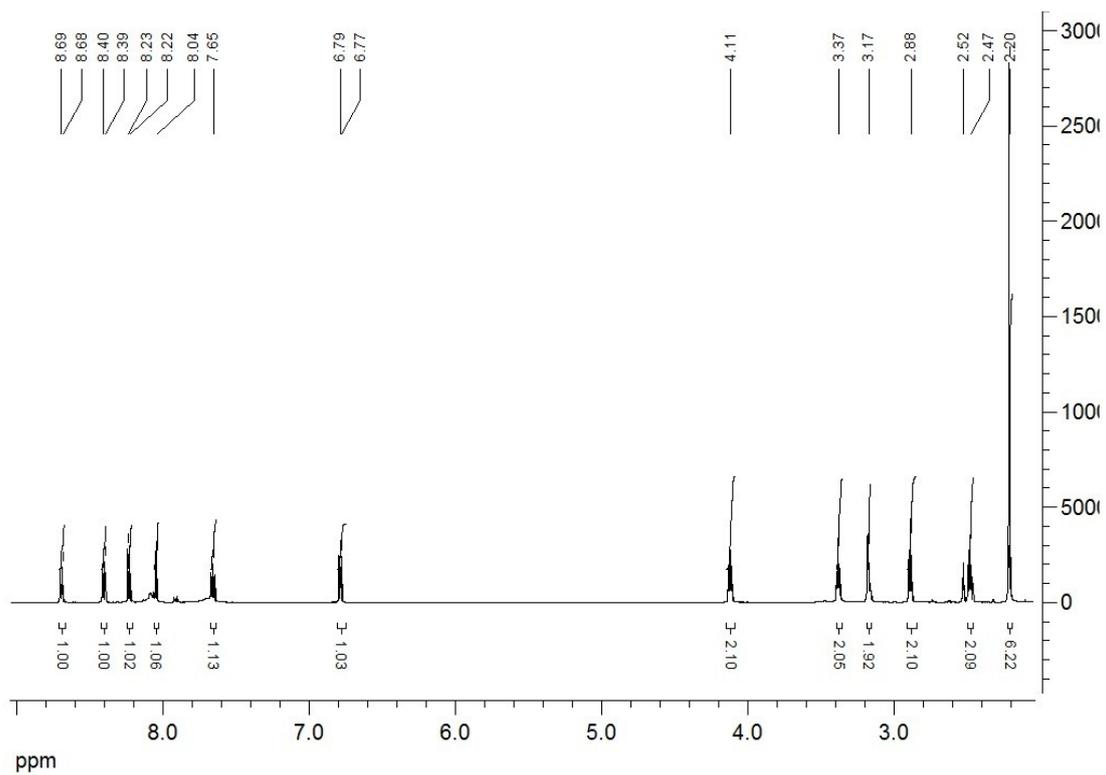


Fig. S8 ^1H NMR of compound **1** in DMSO-d_6 .

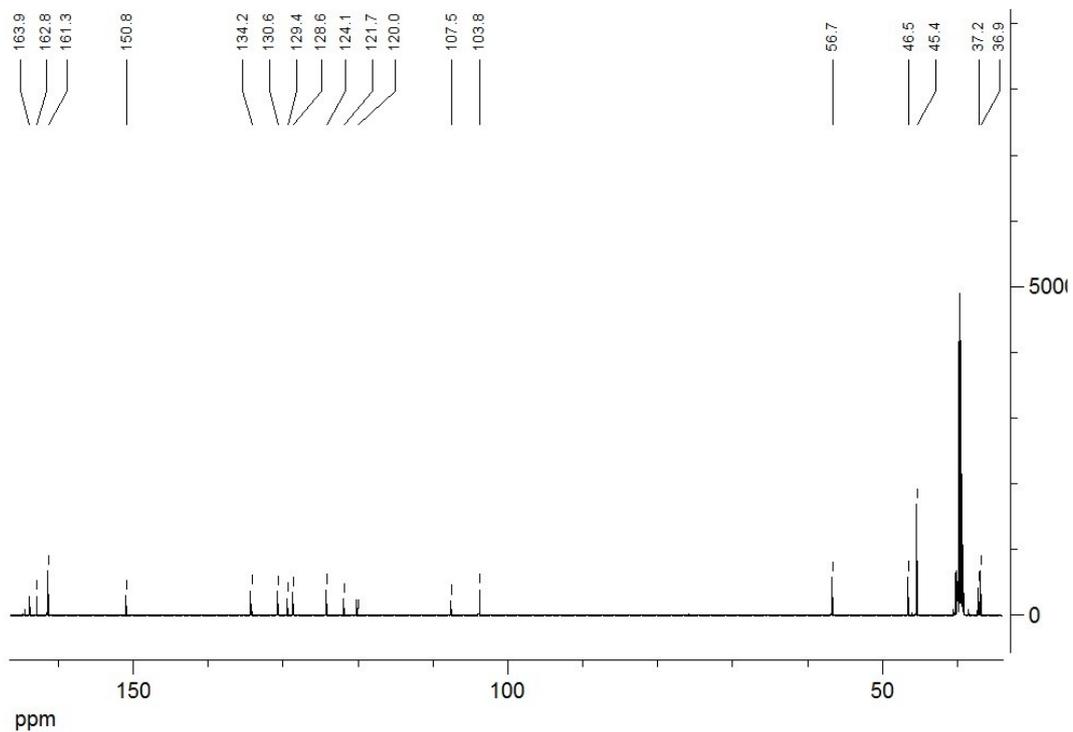


Fig. S9 ^{13}C NMR of compound **1** in DMSO-d_6 .

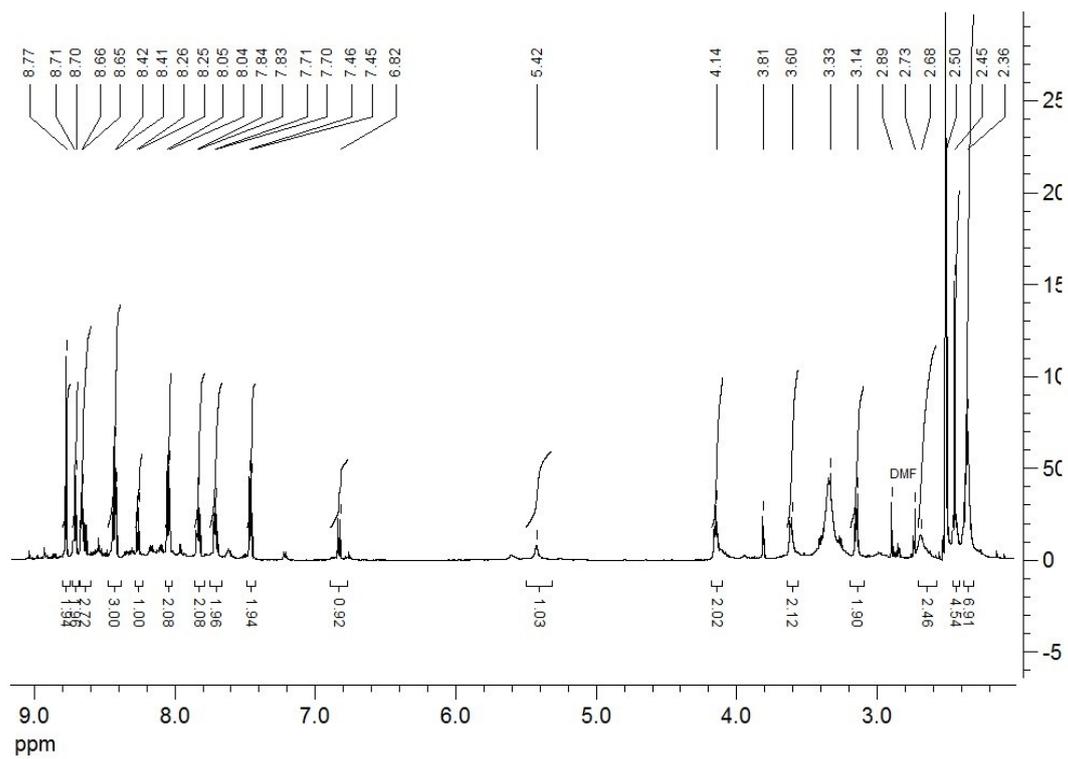


Fig. S12 ^1H NMR of complex **3** in DMSO-d_6 .

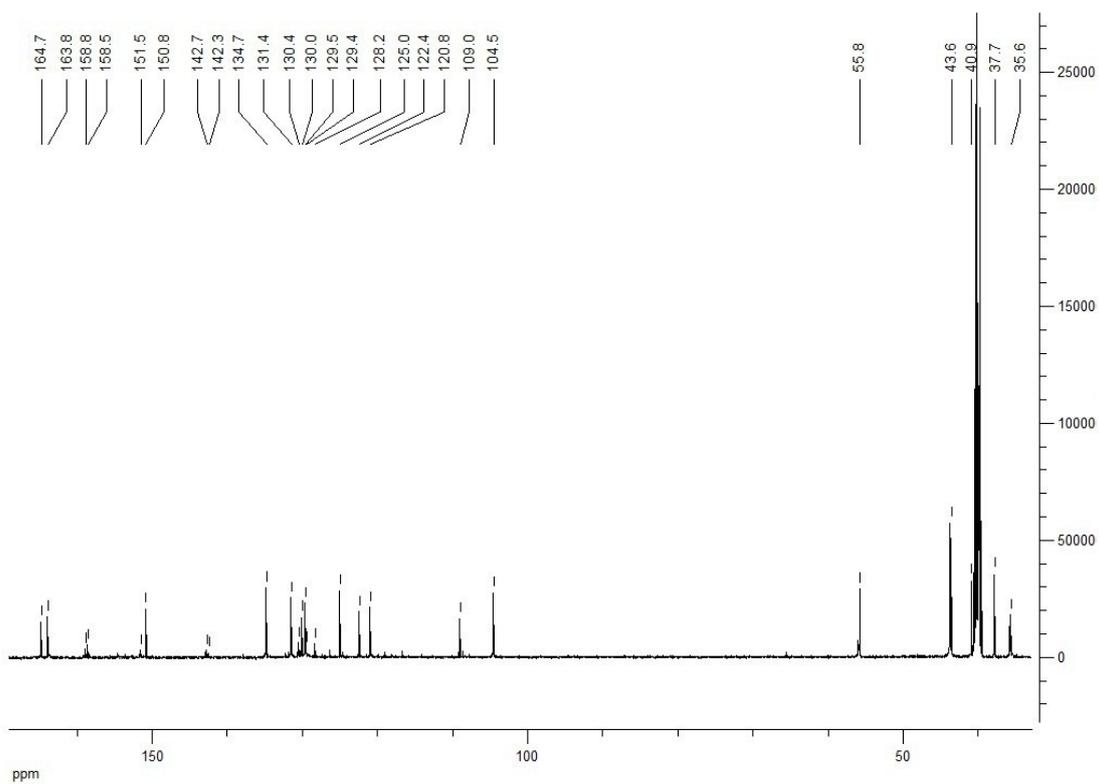


Fig. S13 ^{13}C NMR of complex **3** in DMSO-d_6 .