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A Ruthenium(II) complex as a potential luminescent switch-on probe for G-quadruplex DNA†

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This study aimed to synthesize two ruthenium(II) complexes modified with aryl alkynes [Ru(phen)₂(TMSEPIP)](ClO₄)₂ (**1**, phen = 1,10-phenanthroline; TMSEPIP = 2-(2-trimethylsilylethylphenyl)imidazole[4,5*f*][1,10]phenanthroline) and [Ru(phen)₂(BEPIP)](ClO₄)₂ (**2**, BEPIP = 2-(2-phenethylphenyl)imidazole[4,5*f*][1,10]phenanthroline) and evaluate their ability to act as luminescent probes of *c-myc* G-quadruplex DNA. It was discovered that the luminescence of **1** decreased in the presence of *c-myc* G-quadruplex DNA, while the luminescence of **2** clearly increased under the same conditions. Besides, in the presence of double-stranded DNA, the luminescence intensity of **1** and **2** showed little obvious change. It was interesting to find by atomic force microscopy that **1** could promote *c-myc* G-quadruplex DNA self-assembly to form a nano-network structure, while **2** could promote the formation of a nanowire structure. Further studies showed that **1** was inserted into the groove formed by base pairs A6-G8 and G21-T23 of *c-myc* G-quadruplex DNA, while **2** was inserted into the groove formed by G7-G9 and G22-A24. It is speculated that **2** can be developed as a potential luminescent switch-on probe through selectively recognizing and promoting self-assembly of *c-myc* G-quadruplex DNA.

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1 Introduction

In recent years, much attention has been focused on the self-assembly of the G-quadruplex, a special conformation formed by a G-rich sequence through Hoogsteen hydrogen bonding, for its potential utility as a gene carrier and in targeted drug delivery systems and nanodevices.¹ It has been reported that G-rich oligonucleotides such as d(GCGGAGGCG) can self-assemble to form nanowires, and tetramolecular G-quadruplexes can also assemble to form nanowires when guided by parallel-stranded duplexes.^{2,3} In general, DNA self-assembly can be successfully implemented using DNA tiles, DNA origami, and dynamic self-assembly.⁴ In addition, in recent years, the in-depth understanding of noncovalent interactions of small molecules with biological macromolecules encouraged spontaneous DNA assembly into remarkably diverse architectures using synthetic small molecules.^{5,6} For example, Co(II) complexes have been used to induce rapid DNA condensation into globular nanoparticles through

intermolecular π - π stacking interactions.⁷ Moreover, polypyridyl Ru(II) complexes ([Ru(phen)₂(bqbg)]²⁺) and ([Ru{(bpy)₂-Ru(H₂bpib)}₃]Cl₈) have also facilitated induction of double-strand DNA condensation into nanoparticles/nanowires.⁸ It has been revealed that condensed nanoparticles of DNA formed using this method can be taken up by cells and used as a one- and two-photon luminescent tracking nonviral gene vector.⁹

Nevertheless, the self-assembly of G-quadruplex DNA promoted by inorganic materials, especially Ru(II) complexes, still remains uncovered. Accumulated evidence shows that Ru(II) complexes exhibit excellent binding affinity to G-quadruplex DNA. Chao reported that octahedral polypyridyl Ru(II) complexes could recognize and bind to telemetric G-quadruplex DNA.¹⁰ It has also been revealed that arene Ru(II) complexes can bind and stabilize *c-myc* G-quadruplex in a groove binding mode.¹¹ Further studies also showed that nanosized chiral supramolecular complexes preferentially bind to higher-order G-quadruplexes over single G-quadruplex units with ~200-fold different selectivity, and that dimeric G-quadruplex units can be novel binding sites for large ligands targeting higher-order G-quadruplexes.¹² Recently, it has been reported that specific Ru(II) dppz compounds can distinguish G-quadruplex from duplex DNA. The photophysical process controlling the DNA-sensing properties has been explored; multiple models have been proposed to explain this “light-switch” behavior. So, needs to be clarified whether the Ru(II) complexes can act as a luminescent “switch-on” probe through the high selectivity of G-quadruplex DNA.

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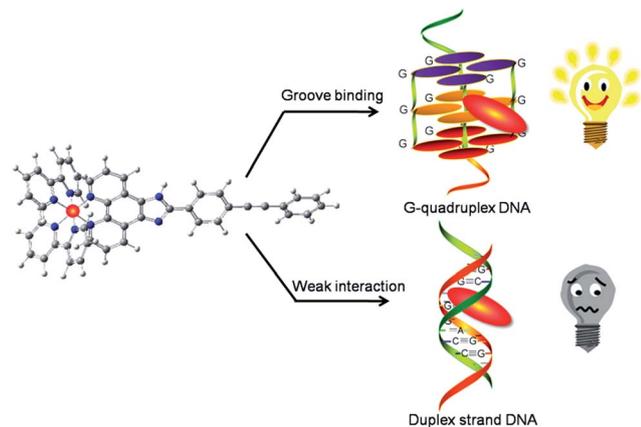
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Scheme 1 A ruthenium(II) complex with phenylethynyl exhibited stronger affinity with G-quadruplex DNA than ds-DNA, which can act as a potential luminescent “switch on” probe for selective recognition of G-quadruplex DNA.

This study reported the preparation of two ruthenium(II) complexes modified by aryl alkyne group using Sonogashira C–C coupling reaction under microwave irradiation (see ESI Fig. S1†), and their interactions with *c-myc* G-quadruplex DNA have also been studied. The results showed that both complexes exhibited stronger affinity to G-quadruplex DNA than to double-strand DNA. However, they exhibited different binding sites with *c-myc* G-quadruplex DNA in groove binding mode to promote different self-assembled nanostructures. Moreover, the fluorescence emission of 2 enhanced with the addition of *c-myc* G-quadruplex DNA (Scheme 1). Then, a simple, selective, G-quadruplex-based label-free luminescent switch-on assay was investigated.

2 Experimental section

2.1 Materials and methods

All the chemicals including solvents were obtained from commercial vendors and used as received. Ruthenium(III) chloride hydrate was purchased from Mitsuwa Chemicals. *c-myc* G-quadruplex DNA (5'-TGGGGAGGGTGGGGAGGG TGGGGAAGG-3') and the fluorescent labeled oligonucleotide, *c-myc* G-quadruplex DNA (5'-FAM-TGGGGAGGGTGGGGAGGGTGGGGAAGG-TAMRA-3', FAM: carboxyfluorescein, TAMRA: 6-carboxytetramethylrhodamine) and formed a G-quadruplex conformation by renaturation at 4 °C for 24 h after 90 °C denaturation for 5 min, according to the literature.^{13,14} And *c-myc*, 5'-FAM-*c-myc*-TAMRA-3', ds26/ss DNA and CT-DNA were purchased from Sangon Biotech (shanghai) Co., Ltd. All aqueous solutions were prepared with doubly distilled water. The Tris-HCl buffer consisting of Tris (10 mM) and KCl/NaCl (100 mM), adjust pH value to 7.2 by HCl (0.1 mol), which was applied to UV titrations, fluorescence emission titrations, AFM observation and analytical ultracentrifugation assay. MDA-MB-231 cells were purchased from ATCC, which were cultured by DMEM (HyClone) with 10% fetal bovine serum (HyClone).

Microwave assisted synthesis was carried out on Anton Paar Monowave 300 microwave reactor. The electronic spectra were

recorded on a Shimadzu UV-2550 Spectrophotometer and steady-state emission spectra were recorded on a RF-5301 Fluorescence Spectrophotometer. The ¹H NMR, ¹H-¹H COSY and ¹³C NMR spectra were recorded in DMSO-*d*⁶ solution on a Bruker DRX 2500 spectrometer, and ESI-MS spectra were obtained in methanol on Agilent 1100 ESI-MS system operating at room temperature. The nanostructures were characterized by atomic force microscopy (AFM) (Bruker, Dimension FastScanTM).

2.2 Synthesis and characterization

2.2.1 Synthesis of [Ru(phen)₂(*p*-TMSEPIP)](ClO₄)₂ (1). 1 was synthesized following the literature but with some modifications. In general, [Ru(phen)₂(*p*-BrPIP)](ClO₄)₂ (130 mg, 0.125 mmol) and trimethylsilylacetylene (0.09 mL, 0.625 mmol) was dissolved in dry CH₃CN (15.0 mL), and then Pd(PPh₃)₂Cl₂, CuI and dry Et₃N were added into the reaction system under N₂ atmosphere. The reaction mixture was irradiated by microwaves for 30 min at 140 °C. After filtrated and dried in vacuums, a dark red powder was obtained. The crude was purified by flash silica gel column chromatography (CH₃CN), yields: 30%. ESI-MS (in CH₃CN, *m/z*): 427.4 ([M-2ClO₄]²⁺), 853.3 ([M-2ClO₄-H]⁺). ¹H NMR (400 MHz, DMSO-*d*⁶) δ 9.03 (d, *J* = 8.1, 2H, H_c), 8.79 (t, 2H, H_{4'}), 8.77 (t, 2H, H₄), 8.40 (s, 4H, H_{5,6}), 8.37 (d, *J* = 8.4, 2H, H_e), 8.11 (m, 4H, H_{2'}, H₂), 7.91 (d, *J* = 4.7 Hz, 2H, H_a), 7.78 (ddd, *J* = 8.2, 5.3, 1.8 Hz, 4H, H_{3'}, H₃), 7.73 (dd, *J* = 8.3, 5.3 Hz, 2H, H_b), 7.67 (d, *J* = 8.3 Hz, 2H, H_d), 1.76 (s, 9H, Si(CH₃)₃). ¹³C NMR (101 MHz, DMSO-*d*⁶) δ 171.91 (s), 153.10 (s), 147.73 (d, *J* = 9.6 Hz), 137.16 (s), 132.67 (s), 130.90 (d, *J* = 2.8 Hz), 128.50 (s), 126.88 (d, *J* = 20.1 Hz), 40.66 (s), 40.45 (s), 40.24 (s), 40.04 (s), 39.83 (s), 39.62 (s), 39.41 (s), 22.97 (s).

2.2.2 Synthesis of [Ru(phen)₂(*p*-BEPIP)](ClO₄)₂ (2). 2 was synthesized as the same methods as above, but with phenylacetylene (0.07 mL, 0.625 mmol), yields: 51%. ESI-MS (in CH₃CN, *m/z*): 429.5 ([M-2ClO₄]²⁺), 857.3 ([M-2ClO₄-H]⁺). ¹H NMR (400 MHz, DMSO-*d*⁶) δ 9.04 (d, *J* = 8.1 Hz, 2H, H_c), 8.79 (t, 2H, H_{4'}), 8.77 (t, 2H, H₄), 8.44 (d, *J* = 8.4, 2H, H_e), 8.40 (s, 4H, H_{5,6}), 8.12 (ddd, *J* = 10.2, 5.3, 1.2 Hz, 4H, H_{2'}, H₂), 7.88 (d, *J* = 4.5 Hz, 2H, H_a), 7.78 (ddd, *J* = 8.5, 5.3, 3.5 Hz, 4H, H_{3'}, H₃), 7.75–7.68, (m, 4H, H_b, H_d), 7.61–7.57 (m, 2H, H_g), 7.46 (d, *J* = 2.1 Hz, 1H, H_h), 7.45 (d, *J* = 1.9 Hz, 2H, H_f). ¹³C NMR (101 MHz, DMSO-*d*⁶) δ 153.08 (s), 147.76 (d, *J* = 10.0 Hz), 137.12 (s), 132.24 (s), 131.84 (s), 130.88 (s), 129.29 (s), 128.53 (s), 127.04 (s), 126.78 (s), 125.91 (s), 40.59 (d, *J* = 15.5 Hz), 40.37 (d, *J* = 17.9 Hz), 40.16 (d, *J* = 18.2 Hz), 40.04 (s), 39.75 (d, *J* = 16.5 Hz), 39.53 (d, *J* = 18.3 Hz), 39.41 (s), 22.97 (s).

2.3 Fluorescence properties assay

The fluorescence quantum yields Φ (relative values) of samples were calculated according to the following expression:¹⁵

$$\Phi_{\text{unk}} = \Phi_{\text{std}} \left(\frac{I_{\text{unk}}}{A_{\text{unk}}} \right) \left(\frac{A_{\text{std}}}{I_{\text{std}}} \right) \left(\frac{\eta_{\text{unk}}}{\eta_{\text{std}}} \right)^2$$

where the subscripts unk and std respectively refer to the sample and the reference, Φ_{unk} and Φ_{std} are the fluorescence quantum yields of the sample and standard, respectively. I_{unk} and I_{std} are the integrated emission intensities of the sample



and standard, respectively. A_{unk} and A_{std} are the absorbances of the sample and standard at the best excitation wavelengths of sample, respectively, and η_{unk} and η_{std} are the indexes of refraction of the sample and standard solutions. The fluorescence quantum yields of two complexes were measured at best excitation wavelengths using $\text{Ru}(\text{bpy})_3^{2+}$ in CH_3CN as the radiative quantum yield standard. Fluorescence spectroscopy measurements were performed using a 1 cm path length quartz cell. Luminescence titrations of $\text{Ru}(\text{II})$ complexes were measured by using fixed ruthenium concentration (4 μM) with increasing *c-myc* G4 DNA or CT-DNA. The first spectrum was taken after 5 min to allow the sample (3 mL) to equilibrate. Then, 2 μL of a 100 μM DNA solution was added to the sample cell with thorough mixing. The titration processes were repeated until there was no apparent.

2.4 Atom force microscopy observation

The mixed solution of DNA (100 μM) and $\text{Ru}(\text{II})$ complexes (100 μM) were incubated for three days. Then, the mixed solution of 10 μL was removed to a mica plate and naturally volatilized for 2 h. An image of the sample was captured by atomic force microscopy (AFM).

2.5 Analytical ultracentrifugation assay

Analytical ultracentrifugation was operated by ProteomeLab XL-A/XL-I (Beckman Coulter). Sedimentation velocity experiments were carried out in a Beckman XL-I analytical ultracentrifuge. Samples were prepared with a final concentration of 100 μM *c-myc* G-quadruplex DNA, *c-myc* G-quadruplex DNA (100 μM) bound 1 and 2 (100 μM) in buffer (Tris-HCl buffer pH 7.2, 10 mM Tris and 100 mM KCl). *c-myc* G-quadruplex DNA and 1 or 2 were mixed 1 : 1 with a final volume of 400 μL . Samples were loaded into a two-channel ultracentrifuge cell, and a blank buffer consisting of the system storage buffers, Tris-HCl buffer was loaded into the blank channel. For interference experiments, sapphire windows were used, and for absorbance experiments, quartz windows were used. Cells were loaded in an An-60Ti rotor and centrifuged at 50 000 rpm at 20 °C. Data were analyzed using SEDFIT using a continuous $c(s)$ with bimodal f/f_0 .¹⁶ Fits were considered satisfactory if the root mean square deviation was less than 0.009 and the residuals were randomly distributed.

2.6 Molecular docking

Automated docking studies were performed with three different docking algorithms, which were: (1) AutoDock 3.0 ('Lamarckian' genetic algorithm); (2) FlexX 1.10 (incremental construction algorithm, as implemented in Sybyl 6.8), and (3) GOLD 1.2 (*i.e.*, the "Darwinian" genetic algorithm). As scoring is a very important second aspect of automated docking methodologies, it was decided to investigate the effect of rescoring – this is a process of reprioritization of the docking solutions (*i.e.*, primarily ranked by the "native" scoring function implemented in the docking program) with an additional stand-alone scoring function.

2.7 Spectroscopy analysis

The recording of electronic absorption spectra was carried out at room temperature to determine the binding affinity between *c-myc* G4-DNA/CT-DNA and ruthenium(II) complexes ($[\textit{c-myc} \text{ G4-DNA}] = 100 \mu\text{M}$ $[\text{CT-DNA}] = 6 \text{ mM}$). 3.0 mL solutions of the blank buffer and the ruthenium complex samples (10 μM) were placed in the reference and sample cuvettes respectively, following which, the first spectrum was recorded in the wavelength range of 200–800 nm. During the titration, an aliquot (2 μL) of buffered DNA solution was added to each cuvette to eliminate the absorbance of *c-myc* DNA/CT-DNA itself, and the solutions were mixed by repeated inversions. After the solutions were mixed for approximately five minutes, the absorption spectra were recorded. The titration process was repeated until there was no change in the spectra for at least four titrations, which indicated that binding saturation had been achieved.

2.8 Melting and competitive FRET assays

The fluorescent labeled oligonucleotide, *c-myc* G-quadruplex DNA (5'-FAM-TGGGGAGGGTGGGGAGGG TGGGGAAGG-TAMRA-3', FAM: carboxyfluorescein, TAMRA: 6-carboxy tetramethylrhodamine) used as the FRET probe was diluted in Tris-HCl buffer and then annealed by being heated to 90 °C for 5 min, followed by slowly cooling to room temperature.¹⁷ Ds26 duplex DNA (CAATCGGATCGAATTCGATCCGATTG) and single-stranded (ss) DNA f(5'-CTCAT4C2ATACAT2A3GATAGTCAT-3') were competitive binders to evaluate the selective binding ability of $\text{Ru}(\text{II})$ complexes with *c-myc* G4 DNA. Fluorescence melting curves were determined with a Bio-RadiQ5 realtime PCR detection system, by using a total reaction volume of 25 mL, with labeled oligonucleotide (1 μM) and different concentrations of complexes in Tris-HCl buffer.¹⁸ A constant temperature was maintained for 30 s prior to each reading to ensure a stable value. Final analysis of the data was carried out by using Origin7.5 (Origin Lab Corp.).

2.9 Western blot analysis

Total cellular proteins were extracted by incubating cells in lysis buffer obtained from Cell Signaling Technology and protein concentrations were determined by BCA assay. SDS-PAGE was done in 10% tricine gels loading equal amount of proteins per lane. After electrophoresis, separated proteins were transferred to nitrocellulose membrane and blocked with 5% non-fat milk in TBST buffer for 1 h. After then, the membranes were incubated with primary antibodies at 1 : 1000 dilutions in 5% non-fat milk overnight at 4 °C, and then secondary antibodies conjugated with horseradish peroxidase at 1 : 2000 dilution for 1 h at room temperature. Protein bands were visualized on X-ray film using an enhanced chemiluminescence system (Kodak).

3 Results and discussion

3.1 The chemical properties of ruthenium(II) complexes

Excitation of 1 (4 μM) at about 462 nm in Tris-HCl-KCl buffer resulted in an emission with λ_{max} at 583 nm, while the excitation of 2 (4 μM) at about 467 nm led to the emission with λ_{max} at



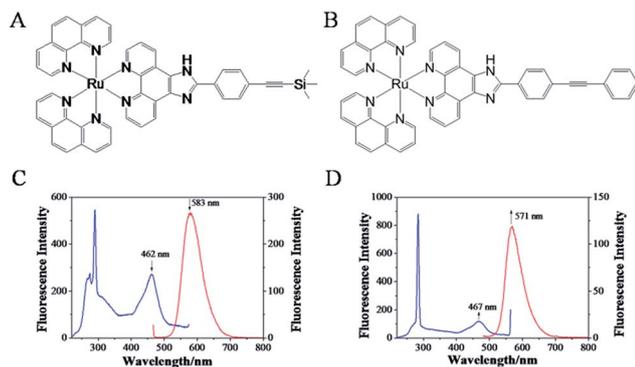


Fig. 1 The synthetic ruthenium(II) complexes with phenylethynyl **1** (A) and **2** (B). The fluorescence properties of **1** (C) and **2** (D). [Ru] = 4 μ M, excitation spectra (blue) and emission spectra (red).

571 nm. The Stokes shift was determined from the excitation and emission spectral peak energies, the noticeable differences between the excitation and emission wavelengths was calculated to be almost 121 nm of **1** and 104 nm of **2**,¹⁹ (Fig. 1) which might contributed to the presence of trimethylsilylacetylene exhibited better electron-donating conjugation effect than phenylethynyl.^{20,21} The quantum yields of **1** and **2** in CH₃CN were 0.028 and 0.016, respectively. Interestingly, it was found that, with the addition of *c-myc* G-quadruplex DNA, the fluorescence intensity of **1** decreased gradually, about 35.6%, which may be attributed to the binding of **1** to *c-myc* G-quadruplex DNA (Fig. 2A). However, for **2**, the fluorescence intensity increased dramatically, about 81.6%, which may be attributed to switch-on assay (Fig. 2B).²² Moreover, strong red fluorescence of **1** (4 μ M) in Tris-HCl-KCl buffer solution was observed. When excited at 365 nm of ultraviolet (UV) light, with the addition of *c-myc* DNA, the fluorescence emission decreased (Fig. 2E); but for **2**, the fluorescence emission increased (Fig. 2F). However, with the addition of CT-DNA, the fluorescence intensity of **1** and **2**

were both decreased slightly (Fig. 2C and D), and the fluorescence emission of **1** and **2** have little obvious change (Fig. 2G and H), which indicated that both ruthenium(II) complexes exhibited weak interactions with duplex strand DNA. Above results suggested that the binding of **2** to *c-myc* G-quadruplex DNA might act as a switch-on luminescent probe to recognize *c-myc* G-quadruplex DNA.^{23–25}

3.2 High-order self-assembly of G-quadruplex DNA controlled by ruthenium(II) complexes

The self-assembly of *c-myc* G-quadruplex DNA induced by both Ru(II) complexes was first confirmed by AFM. Incubation of **1** with *c-myc* G-quadruplex DNA led to the formation of a mesh hole-net structure with diameters of approximately 500–1000 nm (Fig. 3A),^{7,26} which indicated that *c-myc* G-quadruplex DNA self-assembled into an orderly DNA nano-network structure in the presence of **1**. As for *c-myc* G-quadruplex DNA incubated with **2**, a typical pipeline-like structure of approximately 200 nm was observed under AFM (Fig. 3B), indicating that a nanowire structure was formed from the self-assembly of *c-myc* G-quadruplex DNA promoted by **2**.²⁷ Moreover, when amplified, some hole-net structures with an average size of about 250 nm were observed in the nanowires of DNA, indicating that a mixed self-assembled G-quadruplex DNA formed nano-network and nanowire structures (Fig. 3B).²⁸ The aforementioned results indicated that both complexes could promote DNA self-assembly but form different nanostructures, which might be attributed to different binding abilities.

3.3 Molecular conformation of G-quadruplex DNA induced by ruthenium(II) complexes in solution

The oligomeric state of *c-myc* G-quadruplex DNA in solution was investigated in the presence and absence of **1** and **2** by sedimentation velocity analytical ultracentrifugation. The effects of **1** and **2** were further examined as a function molecular size on

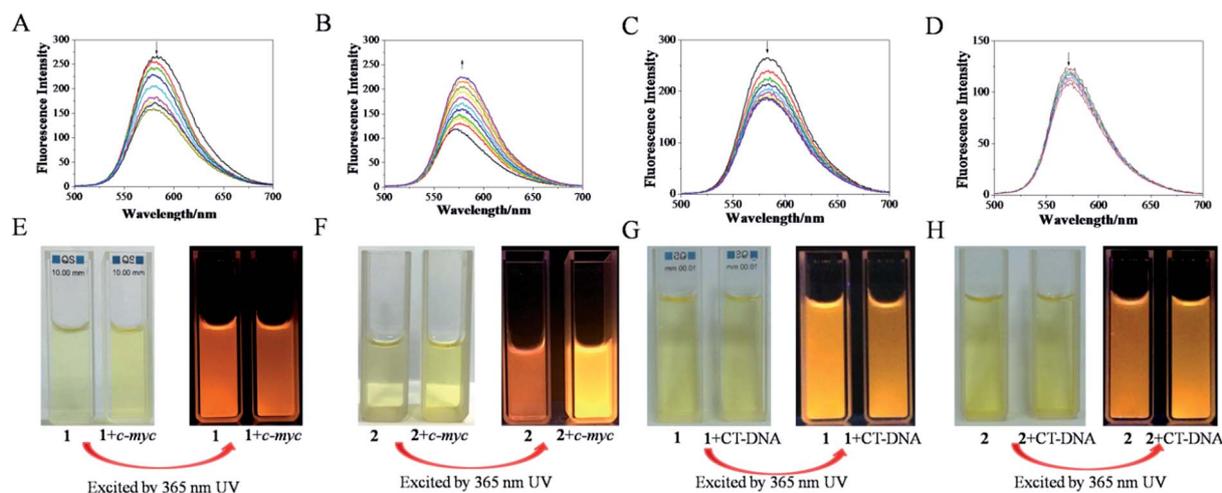


Fig. 2 The changes of fluorescence intensity of **1** (A) and **2** (B) with the addition of *c-myc* G-quadruplex DNA, **1** (C) and **2** (D) with the addition of CT-DNA. ([Ru] = 4 μ M, [*c-myc*] = 0.067*n* μ M, *n* = 0, 1, 2...); the fluorescence emission of **1** (E) and **2** (F) in the absence and presence of *c-myc* G4-DNA in a Tris-HCl-KCl buffer solution (pH 7.2), **1** (G) and **2** (H) in the absence and presence of CT-DNA in Tris-HCl-NaCl buffer solution (pH 7.2), which excited at 365 nm using a portable UV lamp. ([Ru] = 20 μ M, [*c-myc*] = 20 μ M, [CT-DNA] = 20 μ M).



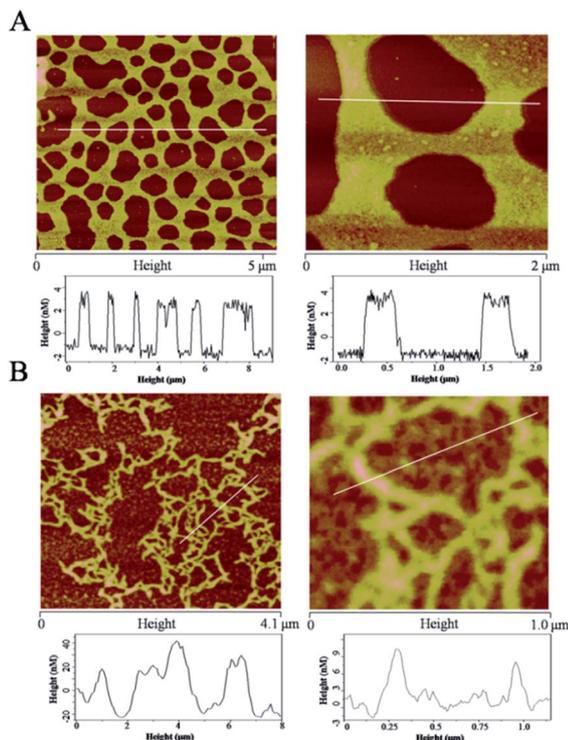


Fig. 3 (A) Atomic force microscopic (AFM) image and analysis of *c-myc* DNA self-assembled into nano-network structures in the presence of **1**; (B) AFM image and analysis of *c-myc* DNA self-assembled into nanowire structures in the presence of **2**. [Ru]:[*c-myc*] = 1 : 1, (100 μ M).

the ability to visually detect molecular conformation change of *c-myc* G-quadruplex DNA. Under higher loading concentration (100 μ M) of free *c-myc* G-quadruplex DNA, the weight average *s*

value shifted slightly to 1.854 S and 3.378 S, and the relative amount of monomer reached to 79.8% and trimer for 18.9% of absorption at the expense of the other species. This was illustrated in Fig. 4d, which suggested that free *c-myc* G-quadruplex DNA in solution displayed two possible state having the following molecular weights, monomer state of 10.6 kDa, trimer state of 26.1 kDa. Then, upon the addition of **1**, the *c-myc* G-quadruplex DNA in solution displayed a good deal of aggregation state with higher than monomer (Fig. 4e), 19.5 kDa and 2.410 S (dimer, 95.091%), 221 kDa and 12.18 S (polymer, 0.132%).^{29,30} However, in the presence of **2**, the sedimentation distributions changed drastically, and 72.13% of the absorbance corresponded to a trimeric species, while the remaining absorbance sedimented as a heterogeneous mixture with *s*-values ranging between 2.09 and 21.4 S, corresponding to highly anisotropic aggregates ranging between 14.5 and 468 kDa in size. It showed that with the increasing of **2**, the *c-myc* G-quadruplex DNA displayed plenty of trimerization state and different aggregation state (Fig. 4f), 29.5 kDa, 2.09 S (trimer), 77.4 kDa, 6.38 S (polymer) and 139 kDa, 9.41 S (polymer). In any case, the sedimentation velocity further revealed that **2** do participate in the association to promote the self-assembly of *c-myc* G-quadruplex DNA to form nanowire.

3.4 Self-assembly orientations of ruthenium(II) complexes with *c-myc* G-quadruplex DNA

To further understand the self-assembly of *c-myc* G-quadruplex DNA enhanced by these Ru(II) complexes, the molecular docking simulations were carried out to confirm the binding G-quadruplex DNA in a typical groove binding mode. Moreover, **1** was inserted into the groove of *c-myc* G-quadruplex DNA,

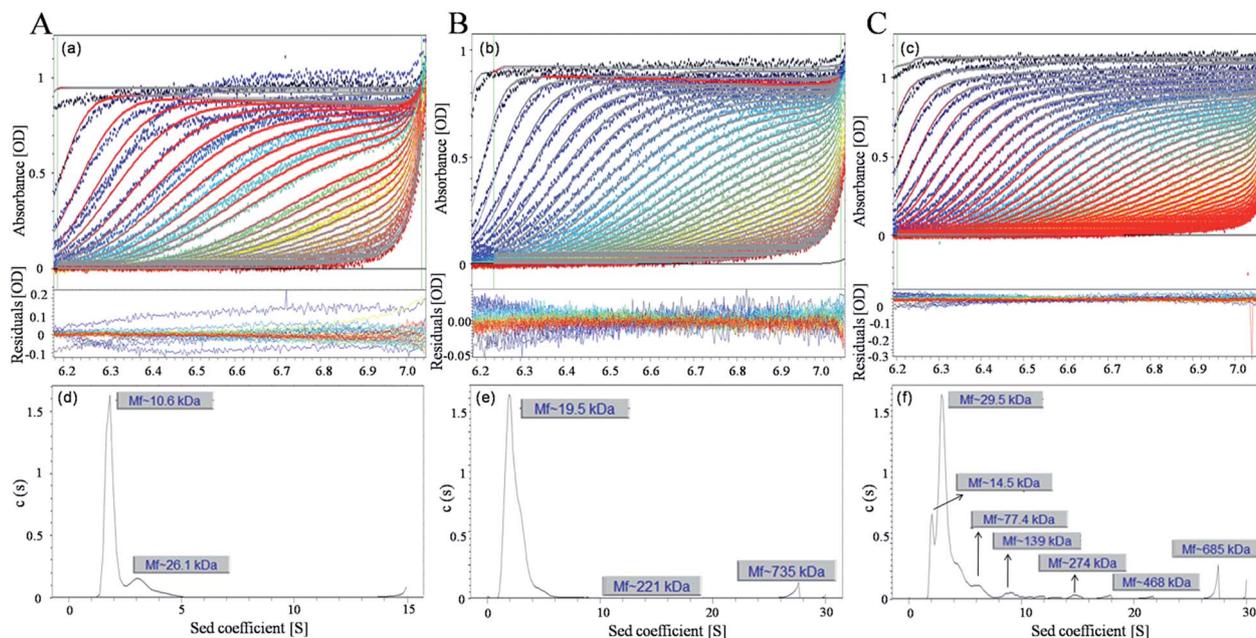


Fig. 4 Analytical ultracentrifugation analyzed the sedimentation velocity (SV) of *c-myc* (A), **1** (B) and **2** (C). All theoretically calculated SV concentration profiles, $s(r, t)$, were generated using SEDFIT, fitting error and fitting curve (a–c). Sedimentation coefficient distributions $c(s)$ from the analysis of the sedimentation profiles of the simulated *c-myc* G-quadruplex-**1/2**-bound DNA system (d–f), SV runs monitored by absorbance at 300 nm. Concentrations are [*c-myc*] = 100 μ M, [**1**] = 100 μ M and [**2**] = 100 μ M.



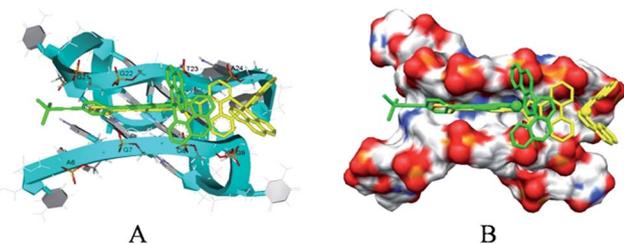


Fig. 5 The binding sites (A) and binding mode (B) of **1** and **2** with *c-myc* G4 DNA calculated by molecular docking.

which was formed by base pairs A6-G8 and G21-T23 with a binding energy of -6.29 kcal mol $^{-1}$. However, **2** was found to insert into the groove of *c-myc* G-quadruplex DNA, which was *c-myc* G-quadruplex DNA led to enhanced fluorescence, formed by base pairs G7-G9 and G22-A24 with a binding energy of -7.83 kcal mol $^{-1}$ (Fig. 5).¹¹ As for **1**, the presence of trimethylsilylacetylene provided steric hindrance, and the molecules were inserted into a shallow groove of G-quadruplex DNA. However, for **2**, the presence of phenylethynyl provided better planarity, and the molecules were inserted into a deep groove of G-quadruplex DNA through π - π stacking. These data indicated that both Ru(II) complexes might bind to *c-myc* G-quadruplex DNA in a groove binding mode, and **2** exhibited stronger binding ability than **1**.¹² Moreover, different binding sites guide different self-assembly nanostructures of *c-myc* G-quadruplex DNA.

3.5 G4/ds DNA-binding properties

Electronic spectra is one of the most common methods to investigate the interactions of small molecules with biological macromolecules. As shown in Fig. 6A, the electronic spectra of **1** exhibited the characteristic intraligand (IL) charge transfer absorption in the range of 250–300 nm with a maximum at 263 nm, upon the addition of *c-myc* G4-DNA, the hypochromism value was 42.3%. For **2**, the hypochromism value was 45.0% (Fig. 6B). However, further analysis upon the addition of CT-DNA showed that both ruthenium(II) complexes bind to CT-DNA in weak affinity (Fig. 6C and D). These data demonstrated that both complexes displayed higher affinity with the *c-myc* G4 DNA than that of CT-DNA; especially **2**, exhibited stronger affinity to *c-myc* G4-DNA.^{31,32}

3.6 Selective recognition of G-quadruplex DNA

To further confirm the selectivity of Ru(II) complexes with *c-myc* G4 DNA, FRET melting assay was carried out to investigate the melting changes of **1** and **2** to *c-myc* G-quadruplex DNA. As shown in Fig. 7, the melting point of *c-myc* G4 DNA was about 41.2 °C, while upon the addition of 6.0 μ M of **1** and **2**, the melting point of *c-myc* G4 DNA (the T_m) increased 46.9 ($\Delta T_m = 5.7$) and 55.9 ($\Delta T_m = 14.7$) °C, respectively. These data indicated that both complexes can stabilize the *c-myc* G4 DNA, especially **2**, displayed better stability than **1**, which accord with aforementioned results. Furthermore, the competitive FRET assay

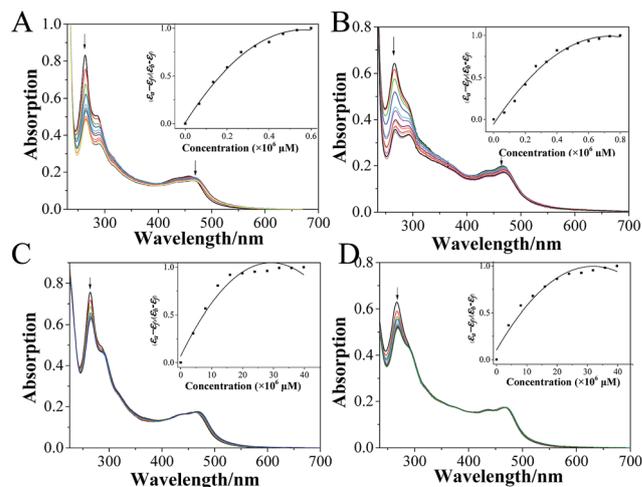


Fig. 6 The study of the interactions between **1** and **2** with *c-myc* G4 DNA/CT-DNA using spectroscopic method. The electronic spectra of **1** (A) and **2** (B) in the absence and presence of *c-myc* G4 DNA in the Tris-HCl-KCl buffer solution (pH 7.2), **1** (C) and **2** (D) in the absence and presence of CT-DNA in the Tris-HCl-NaCl buffer solution (pH 7.2), [Ru] = 10 μ M, [*c-myc* G4 DNA] = 0–0.66 μ M, [CT-DNA] = 0–60 μ M.

was also performed to confirm the selectivity of both Ru(II) complexes between *c-myc* G-quadruplex DNA and single or duplex strand DNA. According to the results of FRET competitive experiments, the melting point for *c-myc* G4 DNA undergo

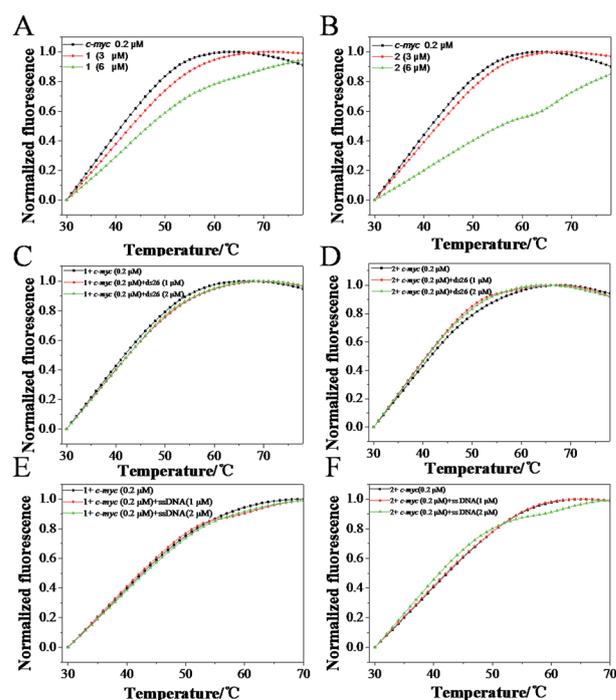


Fig. 7 FRET melting profiles of *c-myc* G4 DNA in the absence and in presence of **1** (A) and **2** (B) ([*c-myc* G4 DNA] = 0.2 μ M) and the competition FRET-melting assay with ds26 in the presence of 3 μ M **1** (C) and **2** (D) and with ssDNA in the presence of 3 μ M **1** (E) and **2** (F), ([Ru] = 3 μ M, [*c-myc* G4 DNA] = 0.2 μ M).



almost little obvious change in the presence of excess duplex ds26 DNA or ssDNA. When $r([\text{ds26}]/[c\text{-myc}]) = 10$, the ΔT_m for **1** and **2** was about 0.86 and -0.9 °C with a drop of less than 5%, respectively (Fig. 7C and D). Moreover, with the excessive supplies of ssDNA, it was also observed that no remarkable change of ΔT_m in the presence of **1** and **2** (Fig. 7E and F). The results indicated that this class of Ru(II) complexes could selectively bind to G-quadruplex DNA more strongly than to single or double-strand DNA, which could act as a switch-on luminescence probe to selective recognize *c-myc* G-quadruplex DNA.^{33–35}

4 Conclusions

In conclusion, two ruthenium(II) complexes modified by aryl ethyl group, $[\text{Ru}(\text{phen})_2(\text{L})](\text{ClO}_4)_2$ (L = TMSEPIP, **1**; BEPIP, **2**) were synthesized in this study using Sonogashira coupling reaction under microwave irradiation. It was confirmed that the luminescence of **1** decreased in the presence of *c-myc* G-quadruplex DNA, the luminescence of **2** increased obviously under the same conditions. It was interesting to find by AFM that **1** could promote *c-myc* G-quadruplex DNA self-assembly to form a nano-network structure, while **2** formed a nanowire structure. Further studies showed that **1** was inserted into the groove formed by base pairs A6-G8 and G21-T23 of *c-myc* G-quadruplex DNA, while **2** was inserted into the groove formed by G7-G9 and G22-A24. It is speculated that **2** can be developed as a potential luminescent switch-on probe through selectively recognizing and promoting self-assembly of *c-myc* G-quadruplex DNA.

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