V-Shaped Dinuclear Pt(II) Complexes: Selective Interaction with Human Telomeric G-quadruplex and Significant Inhibition towards Telomerase

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A quaternized trigeminal ligand, 4-[4,6-di(4-pyridyl)-1,3,5-(2-triazinyl)]-1-methylpyridine-1-ium hexafluorophosphate (dptmpPF6), and two derivative V-shaped dinuclear Pt(II) complexes, \{[Pt(dien)]2(dptmp)](PF6)5 \} (1) and \{[Pt(dpa)]2(dptmp)](PF6)5 \} (2), were synthesized, characterized and applied to a series of biochemical studies. FRET and SPR analyses showed these compounds, especially Pt(II) complexes, bound more strongly to human telomeric (hTel) G-quadruplex than to promoters (such as c-myc and bcl2) or to the duplex DNA. PCR-stop assays revealed that the Pt(II) complexes could bind to and stabilize G-quadruplex far more effectively than corresponding ligand. CD analyses further indicated the three compounds likely stabilized the formation of mixed-type parallel/antiparallel G-quadruplex structures. Their efficacy as telomerase inhibitors and potential anticancer drugs was explored via TRAP. The IC50 value was determined to be 0.113 ± 0.019 μM for 1, indicating that it is one of the strongest known telomerase inhibitors. These results confirm that both V-shaped dinuclear Pt(II) complexes act as selective G-quadruplex binders and significant telomerase inhibitors.

In recent years, G-quadruplexes, which consist of four-stranded nucleic acids, and are described as presenting a high-order secondary DNA structure, have drawn attention regarding their potential use in anticancer therapies1-2. These guanine-rich sequences exist ubiquitously in significant regions of the eukaryotic genome, such as in the promoter regions of several oncogenes and the telomeres at the ends of the chromosomes3-6. It has been reported that telomerase is active and up-regulated in approximately 85% of tumor cells, which would lead to telomere elongation and contribute to cancer cell immortalization7,8. Thus, the telomeric G-quadruplex has been considered to be a potentially effective antitumor target1,2. The formation of a G-quadruplex would result in the inhibition of telomerase activity and would thereby terminate telomere maintenance9. Researches aimed at the stabilization of the G-quadruplex structure of certain sequences and efficiently inhibition of telomerase activity represent a rising field of research in anticancer drug design and development.

G-quadruplexes consist of stacked G-tetrads, which are formed from four guanine bases of G-rich sequences connected by Hoogsteen hydrogen bonds (Fig. 1a). Monovalent cations, such as potassium and sodium ions, can stabilize G-quadruplex structures, presumably via electrostatic interactions with the guanine carbonyl moieties. Previous studies addressing various G-quadruplex conformations have been reported, and crystallographic and NMR data have revealed that G-quadruplexes can be classified according to different strand orientations as showing a parallel structure10 (Fig. 1b), mixed-type “(3 + 1) hybrid” structure11 (Fig. 1c) or basket-type “(2 + 2) hybrid” structure12 (Fig. 1d). These structural data provide important information for the development of G-quadruplex binders.

In past decades, many small molecules that interact with G-quadruplexes have been reported13,14. Some synthetic simple organic molecules15-20, typically metal complexes21-30, have been well characterized as effective G-quadruplex binders. Platinum(II) complexes, as successful inorganic anticancer drugs, have long been considered important due to their interaction with nucleic acids. Recently, planar platinum(II) complexes24-30 have drawn increasing interests related to targeting telomeric G-quadruplex DNA. Some mononuclear platinum(II) complexes, including platinum(II)-phenanthroline24,25, platinum(II)-terpyridine26 and others, with positively
charged side arms, which interact readily with the negatively charged phosphate backbones, grooves and loops of the quadruplexes, have been reported. Additionally, a chain-like bidentate dinuclear platinum(II) complex has been reported as a c-myc and hTel quadruplex binder. A kind of self-assembled tetranuclear platinum(II) complexes with an aromatic plane was studied, which turned out to be good G-quadruplex binders, displaying evident telomerase inhibition. These studies have suggested that a large electron-deficient π-aromatic surface, positively charged area and positively charged substituents represent characteristic features of a good G-quadruplex binder.

We have previously reported a series of mononuclear and tetranuclear Pt(II) complexes as G-quadruplex binders. To further investigate the structure-activity relationship of various Pt(II) complexes, herein we synthesized a new bridged ligand by quaternizing a branch of a trigeminal star-like molecule, 4-(4,6-di(4-pyridyl)-1,3,5-(2-triazinyl))-1-methylpyridine-1-ium iodide (dptmp) with ammonium hexafluorophosphate.

Figure 1 | Schematic representation of a G-tetrad and different conformations of G-quadruplexes; ORTEP views of dptmp and 1. (a) a G-tetrad consists of four guanines, (b) the wild-type 22-mer crystal with K + (parallel structure, PDB ID: 1KF1), (c) the modified 20-mer in K + solution (mixed-type ‘(3 + 1) hybrid’ structure, PDB ID: 2KZD), (d) the wild-type 22-mer in Na + solution (basket-type ‘(2 + 2) hybrid’ structure, PDB ID: 143D).

Results

Synthesis of dptmp and Pt(II) Complexes \([\text{Pt(L^1)}_2\text{(dptmp)}](\text{PF}_6)_3\.

The compound dptmp·1 precipitated from trichloromethane and could not undergo further substitution reactions. After heating the reaction mixture under reflux in trichloromethane for 48 h, it was filtered, the solvent was removed under reduced pressure and the residue was washed with trichloromethane. The isolated dptmp·1 was characterized by H and C NMR spectroscopy, mass spectrometry and elemental analyses (see Experimental Details). Subsequently, dptmp·PF6 was obtained through the reaction of dptmp·1 with ammonium hexafluorophosphate.

The traditional method for synthesizing \([\text{Pt(dien)}\text{Cl}]\text{Cl}-\text{HCl}\) involves the reaction of K2PtCl4 with diethylenetriamine (dien) dissolved in water (pH 3.0) by heating under reflux for 72 h followed by concentration until the yellow product precipitates out. We employed a similar method, described by Annibale et al., using \([\text{Pt(COD)Cl}]_2\) as an intermediate, which was found to be more convenient and resulted in a very high yield of 96.7%.

The complex \([\text{Pt(L^1)}\text{Cl}]\text{Cl}\) was first treated with AgNO3 to eliminate the two chlorides, then filtered to remove AgCl and transferred to a stoppered flask, followed by the addition of dptmp·PF6 and heated at 90°C for 72 h under N2. Finally, excess ammonium hexafluorophosphate was added to the reaction mixture, and the obtained precipitate was filtered out. The two dinuclear Pt(II) complexes were fully characterized via H, C and Pt NMR spectroscopy and elemental analyses. Additionally, dptmp·1 and 1 were structurally characterized by X-ray crystallography (Fig. 1).

Selectively stabilize hTel G-quadruplex DNA structure. Fluorescence resonance energy transfer (FRET) studies were conducted to investigate the binding abilities of the three compounds to human telomeric, promoter G-quadruplex DNA sequences (hTel, c-myc and bcl2) and a duplex DNA sequence. Reliable FRET melting curves (Fig. 2) and stabilization temperature (ΔTm values (Table 1) were obtained.

To quantitatively investigate the binding constants between compounds and DNA samples (the hTel G-quadruplex and duplex DNA sequences), surface plasmon resonance (SPR) experiments were conducted (Supplementary Fig. S8 and Table 2), which serve as a powerful technique for monitoring molecular reactions in real time.

Figure S8 shows the SPR sensorgrams generated for the three compounds binding to the immobilized hTel G-quadruplex and duplex DNA at different concentrations. Based on the obtained data, shown in Table 2, the two Pt(II) complexes displayed higher binding constants with the hTel sequence compared to that with dptmp (4.00 × 105; 1, 1.37 × 105 and 2, 1.40 × 105). Regarding the binding selectivities between the hTel quadruplex and duplex sequences, 1 showed a higher selectivity of 7.1 fold (5,170 vs. 731 nM), compared to 2 at 4.7 fold (3,390 vs. 712 nM), while dptmp showed the lowest selectivity of 1.2 fold (2,890 vs. 2,500 nM). We also found that the kinetic binding affinities (Ku) of the three compounds with the hTel G-quadruplex were superior to those with the duplex DNA sequence.

Stabilizing hTel G-quadruplex structures. To further evaluate whether the compounds effectively bound to the tested oligomer HTG21 and stabilized the G-quadruplex structure, polymerase chain reaction (PCR) stop assays were performed. In these assays, in the presence of the compounds, the template sequence can form a G-quadruplex structure, and PCR products of dsDNA will be undetectable.
Figure 2 indicates that the inhibitory effects of complexes 1 and 2 were enhanced distinctly as the concentrations of both complexes increased from 1.0 to 9.0 μM, with no PCR products being detected at 5.0 or 7.0 μM, respectively. However, dptmp showed a weaker ability to inhibit the appearance of the PCR products and some PCR products were detected at 180 μM. The results indicated that the two complexes exhibited higher binding affinities for the G-quadruplex and were effective G-quadruplex binders. Moreover, to avoid any possible interaction of rTaq polymerase with the compounds and subsequent inhibition of the PCR process, we conducted parallel experiments using the mutated oligomer HIG21 mu instead of HIG21 under the same experimental conditions. The HIG21 mu sequence was unable to form a G-quadruplex. The obtained results suggested that the compounds could not significantly affect the activity of rTaq polymerase at comparable concentrations (Supplementary Fig. S10).

**Induction of the formation of mixed-type parallel/antiparallel hTel G-quadruplexes and stabilization of this structure.** Circular dichroism (CD) spectroscopy was performed to roughly characterize
the structural conversion of various conformations of human telomeric G-quadruplexes, which consist of intra- and inter-molecular in parallel, antiparallel-stranded and mixed arrangements, depending on the strand orientations. It has been reported that the CD spectra of the human telomeric DNA sequence (AG22) may consist of a mixed-type parallel/antiparallel G-quadruplex conformation in the presence of potassium cations, and usually show a strong positive peak around 290 nm, a small positive peak at 265 nm and a small negative peak at 240 nm. When different metal cations, the CD spectra consist of a positive band near 290 nm and a negative band at 265 nm, which may be characteristics of a typical antiparallel G-quadruplex structure. In the absence of metal ions, the CD spectra of the AG22 sequence indicate the coexistence of single strand, parallel and antiparallel G-quadruplexes.

In the current studies, the AG22 sequence was used to study the structural induction and structural transition of G-quadruplexes by CD titration assays in the presence of the three compounds. As reported previously, the AG22 sequence is considered to show a summation spectrum involving multiple G-quadruplexes (mixed-type/basket-type) in K⁺ solution from NMR data obtained for AG22. The sequences Tel26 and wild-type Tel26 which both contain the AG22 four-G-tract human telomeric core sequence and have been well-defined based on NMR and CD data obtained with mixed-type G-quadruplexes, exhibit a distinct CD spectrum including a strong positive peak around 295 nm, with a shoulder peak around 268 nm and a smaller negative peak at 240 nm. When different concentrations of the compounds were added to the K⁺ solution of AG22, very similar CD spectra (Fig. 3) were generated to the Tel26 and wild-type Tel26 which had been confirmed the mixed-type G-quadruplex structure. In the absence of any metal cations, the CD spectra consist of a positive band at 257 nm, a small positive band at 295 nm and a small negative peak near 240 nm, as shown in Figure 3. When the concentrations of the compounds were increased, there was an obvious decrease observed at 257 nm, accompanied by a sharp increase around 295 nm. These data suggest that all of the compounds, especially the two Pt(II) complexes, probably induced the formation of a mixed-type parallel/antiparallel G-quadruplex structure and stabilized this structure. These CD phenomena were consistent with the findings of Zhou et al. Similar conclusions were drawn based on a more identifiable conformation transition detected using 100 mM NaCl, and parallel experiments also roughly demonstrated the selectivity of the compounds towards the mixed-stranded topology.

Inhibition of telomerase activity. To obtain further information about whether the compounds could inhibit telomerase activity and affect telomere elongation by telomerase, we performed the telomeric repeat amplification protocol (TRAP) assays, which can qualitatively and quantitatively evaluate the inhibition of telomerase. In these assays, the elongated telomeres were further amplified by PCR cycling and could be visualized with dye staining. From the results presented in Figure 4, it is clear that the three compounds inhibited telomere elongation in a concentration-dependent manner, while the amplification of the internal telomerase assay standard (ITAS) control was not obviously affected. Thus, it was the telomerase activity rather than the PCR procedure itself that was inhibited by the three compounds. The inhibition of telomerase by the two complexes was found to be due to their function as strong G-quadruplex binders, showing IC₅₀ values of 0.113 ± 0.019 and 3.69 ± 0.53 μM, respectively. Thus, complexes 1 and 2 exhibited stronger telomerase inhibition than dptmp (IC₅₀ = 12.3 ± 1.2 μM).

### Table 1 | Stabilization temperatures, ΔTm [°C] for hTel, c-myc, bcl2 and duplex DNA stabilized by the three compounds determined from FRET

| Compounds | hTel | c-myc | bcl2 | duplex | Cm/K/CDNA |
|-----------|------|-------|------|--------|-----------|
| dptmp     | 17   | 4     | 0    | 0      | 1.25      |
| 1         | 35.4 | 3     | 0    | 0      | 1.25      |
| 2         | 30.5 | 6     | 3    | 1      | 1.25      |

*The concentration of hTel, c-myc, bcl2 and duplex DNA was 400 nM, with a 0.5 μM concentration of the compounds in 60 mM potassium cacodylate buffer (pH 7.4).*

### Table 2 | Kinetic parameters determined with SPR spectroscopy

|          | hTel | duplex | 1       | hTel | duplex | 2       | hTel | duplex |
|----------|------|--------|---------|------|--------|---------|------|--------|
| kₐ (M⁻¹ s⁻¹) | 1.40 × 10⁴ | 1.56 × 10⁴ | 1.12 × 10⁵ | 1.83 × 10⁵ | 1.27 × 10⁵ | 3.60 × 10⁴ |
| kₐ (M⁻¹ s⁻¹) | 3.51 × 10⁻² | 4.50 × 10⁻² | 8.22 × 10⁻² | 9.48 × 10⁻¹ | 9.01 × 10⁻² | 1.22 × 10⁻¹ |
| Kₐ (M⁻¹) | 2.50 × 10⁻⁶ | 2.89 × 10⁻⁶ | 7.31 × 10⁻⁷ | 5.17 × 10⁻⁶ | 7.12 × 10⁻⁷ | 3.39 × 10⁻⁶ |
| Kₐ (M⁻¹) | 4.00 × 10⁸ | 3.46 × 10⁸ | 1.37 × 10⁹ | 1.93 × 10⁹ | 1.40 × 10⁶ | 2.95 × 10⁵ |
| Ch2 [RU] | 6.68 | 9.42 | 5.09 | 2.55 | 10.69 | 15.55 |

*ₕ₀ is the association rate constant, while ₖₐ is the dissociation rate constant.  
ₖₐ was calculated through global fitting of the kinetic data obtained for various concentrations of the compounds using a two-state binding model. Kₐ is given by ₖₐ/ₖ₉.  
The Ch2 value is a statistical measure of the closeness of the fit. Ch2 was of the same order of magnitude as the noise in RU. The fitting process was terminated automatically when a minimum value was found for Ch2.*
Figure 3 | CD titration spectra of the human telomeric G-quadruplexes (3.0 μM) induced by the three compounds (from left to right: dptmp, 1 and 2) in 10 mM Tris-HCl, pH 7.4, at room temperature (r = C_{ML}/C_{DNA}). a: In the absence of metal ions; b: In 100 mM NaCl; c: In 100 mM KCl.

Figure 4 | TRAP assay results for the three compounds (dptmp, 1 and 2), showing telomeric ladders produced by PCR amplification of the oligonucleotides generated by the action of telomerase on a TS primer. The lower band is an internal control primer (ITAS). * Each single assay was dependent on 1.0 μL of telomerase extract (200 cells), prepared from HeLa cells with NP-40 lysis buffer. Negative controls 1 and 2, which accompanied every assay, involved either incubating approximately 1.0 μL of cell lysate at 85°C for 10 min before primer extension or incubating approximately 1.0 μL of NP-40 lysis buffer, respectively. The control (0 nM) assay was performed without adding any compound.
had the highest selectivity. Furthermore, both Pt(II) complexes show preferential quadruplex/duplex DNA selectivity compared to corresponding dptmp, highlighting the importance of the platinum(II) moiety in defining their binding ability. This might be attributed to the platinum(II) moiety that can interact with both the grooves and loops of the hTel quadruplex and their negatively charged phosphate backbones.

The data obtained from PCR stop assays were consistent with previous results, demonstrating that both Pt(II) complexes 1 and 2, especially 1, exhibited an excellent stability towards hTel G-quadruplex, binding more effectively than the corresponding ligand-dptmp. We were able to obtain the exact conformation of the mixed-type quadruplex/duplex DNA structure induced by the two Pt(II) complexes, which is considered to be important for the inhibition of telomerase activity based on the TRAP studies.

In conclusion, we have prepared a water-soluble, quaternized trinuclear ligand and its two derivative Pt(II) complexes. X-ray diffraction analysis showed that the dinuclear Pt(II) complex exhibited a W-shaped structure. Further biochemical assays confirmed that the ligand and the two complexes, especially complex 1, coordinated by straight-chain polyamine, can selectively stabilize the human telomeric G-quadruplex, likely inducing the formation of mixed-type parallel/antiparallel G-quadruplex structure and significantly inhibit the activity of telomerase, probably due to interactions with the sugar-phosphate backbone of the G-quadruplex. Thus, the Pt(II) complexes exhibit a potential promise for antitumor therapies and need to be further investigated to clarify their specific biochemical details.

**Methods**

**Materials.** All chemicals and solvents were obtained from commercial sources and used without further purification.

Synthesis of 4,4′-di-(4-pyridyl)-1,3,5-(2-triazinyl)-1-methylpyridine-1-iodide C30H18N6I (1): A mixture of 4,4′-dipyridyl (0.32 g, 2.00 mmol) and MeI (0.858 g, 6.00 mL) in trichloromethane (240 mL) was stirred at 61°C for 48 h in darkness. The solution gradually became red and a red precipitate formed. The resulting slurry was collected by filtration, and the residue was washed with trichloromethane (2 × 10 mL) to remove tpt. The obtained product was dried under vacuum to give a red solid in 93% yield. 3H NMR (Supplementary Fig. S2) (CD3OD): δ 7.45 (d, J = 4.6 Hz, 4H), 8.98 (d, J = 6.0 Hz, 4H), 7.37 (d, J = 6.0 Hz, 4H), 7.15 (d, J = 8.4 Hz, 4H), 5.01e (s, 3H). 13C NMR (Supplementary Fig. S5) (75 MHz, CD3OD): δ 174.1, 170.8, 154.6, 142.2, 126.7, 126.0, 123.9, 59.4, 58.9, 58.8 ppm, and K2PtCl4 was used as an internal reference (δ = 0). Elemental analysis (analytical, found for [Pt(dien)2(dptmp)PF6]·CH2Cl2: C (30.2, 30.1%), H (2.68, 2.57%), N (14.0, 14.1%), Pt (27.7, 27.8%).

**Fluorescence energy transfer (FRET) studies.** The employed FRET probes included a fluorescein labeled hTel oligonucleotide (5′-FAM-[GGGTTAGGTTAGGTAGTTAGG]-TA9MA-3′, mimicking the human telomeric repeat, TAMRA: 6-carboxytetramethylrhodamine, Sangon), three promoter sequences (c-myc: 5′-FAM-[TGAGGTTAGGTTAGGGTTAGG]-TA9MA-3′, bi-hc: 5′-FAM-[AGGGGCGGCGCGGGGAGGGGGGGAGGGGGGGGCGGCG]-TA9MA-3′, Sangon) and a duplex DNA sequence (5′-FAM-[TATATCCTA-HEG-TATATGTA]-TA9MA-3′, HEG linker: [5′-CH2-CH₂-O]-3′, Sangon). The experiment procedure was as previously reported method. Final analysis of the data was carried out using Origin 8.0 (OriginLab Corp.).

**Surface plasmon resonance (SPR) studies.** SPR measurements were conducted in a ProteOn XPR36 Protein Interaction Array system (Bio-Rad Laboratories, Hercules, CA, USA) using a Neutravidin-coated GLM sensor chip. Biosensor experiments were conducted in filtered and degassed running buffer (Tris-HCl 50 mM pH 7.4, 300 mM KCl, 0.005% Tween-20) at 25°C. The biotinylated oligonucleotides used in these assays were: Biotin-hTel (Quadruplex: 5′-biotin-[AGGGTTAGGTTAGGGTTAGG]-TA9MA-3′; HEG-5′-[FAM-AATGAACTCTGCGGGAATTCG]-TA9MA-3′, Sangon) and DNA sequences were captured (approximately 1,200 RU) in flow cells dptmp-1, 2 and 3, leaving the fourth flow cell as a blank. Data were analyzed with ProteOn manager software, using the two states model for fitting kinetic data.

**Polymerase chain reaction (PCR) stop assays.** The oligonucleotide sequence (HTG21, 5′-[AGGTTAGGTTAGGTAGTTAGG]-3′, Sangon) and the corresponding complementary sequence (HTG21rev, 5′-[ATTCGGCTTCCTGCTCCAA]-3′, Sangon) were used in this assay. The experiment procedure was as previously described method. A parallel experiment was performed using a mutant oligomer, AGG22 (5′-[AGGTTAGGTTAGGTAGTTAGG]-3′, Sangon) with its corresponding complementary sequence HTG21murev (5′-[ATTCGGCTCTGCTAACAA]-3′, Sangon) instead of HTG21 and HTG21rev under the same conditions.

**Circular dichroism (CD) measurements.** CD spectra were measured using a J-810 spectropolarimeter (JASCO, Japan) with a 1 cm cell length quartz cell, over a wavelength range of 220-360 nm at a scan speed of 200 nm/min with five acquisitions at room temperature. The AG22 oligomer (5′-[AGGTTAGGTTAGGTAGTTAGG]-3′, Sangon) was resuspended in Tris-HCl buffer (10 mM, pH 7.4) containing 100 mM KCl, 100 mM Na+ or sodium metatungstate. The experiment procedure was as previously reported method, and the analysis of the data was carried out using Origin 8.0 (OriginLab Corp.).
15 mM MgCl₂, 630 mM KCl, 0.5% Tween-20 and 10 mM EGTA in DEPC-treated 
5 L of dNTP mix (2.5 mM, TaKaRa), 0.4 μL of TS primer (100 ng/μL, 5′-[AACCCCGAAGGAGCTTT]-3′, Invitrogen), 0.8 μL Primer mix (ACX reverse primer, 100 ng/μL, 5′-[GGCGCCGTCCTCCTACCTCACTC]-3′ and NT primer 100 ng/μL, 5′-[ATGCCTGAGGCCTCCTCGTTTT]-3′, Invitrogen), 2.0 μL TSNT internal control primer (4.0 × 10⁻⁵ M, 5′-[AATCCCGAAGGAGCTTTAAGAACGGAGG]-3′, Invitrogen), 0.16 μL of Taq polymerase (5 U/μL, TaKaRa), 5.0 μL of different concentrations of the compounds and 6.64 μL of DEPC-treated 
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**Author contributions**

C.X.X. and Z.W.M. conceived and designed the experiments. C.X.X., Y.X.Z. and X.H.Z. conducted the synthetic experiments. Q.H. conducted the TRAP assays. Y.Z. helped C.X.X. in the TRAP data analysis. L.N.J. provided effective directions during the research process. C.X.X. and Z.W.M. wrote the manuscript. All authors discussed the results and commented on the manuscript. All authors reviewed the manuscript.

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