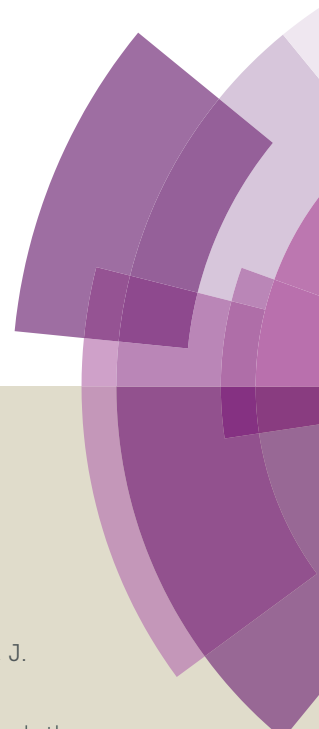
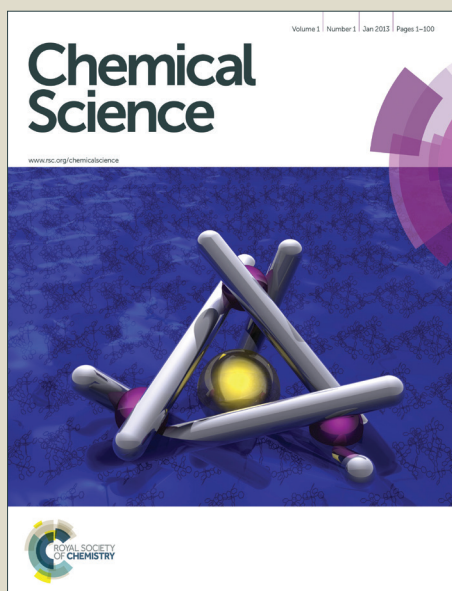


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Terpyridine-Cu(II) Targeting Human Telomeric DNA to Produce Highly Stereospecific G-quadruplex DNA Metalloenzyme†

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The cofactors commonly involved in natural enzymes have provided the inspiration for numerous advances in the creation of artificial metalloenzymes. Nevertheless, to design an appropriate cofactor for a given biomolecular scaffold or vice versa remains a challenge in developing efficient catalysts in biochemistry. Herein, we extend the idea of G-quadruplex-targeting anticancer drug design to construct a G-quadruplex DNA metalloenzyme. We found that a family of terpyridine-Cu(II) complex can serve as an excellent cofactor to dock with human telomeric G-quadruplex DNA. The resulting G-quadruplex DNA metalloenzyme catalyzes an enantioselective Diels-Alder reaction with enantioselectivity over 99% enantiomeric excess and about 73-fold rate acceleration compared to terpyridine-Cu(II) complex alone. The terpyridine-Cu(II) complex cofactor demonstrates dual functions, both as an active site to perform catalysis and as a structural regulator to promote the folding of human telomeric G-quadruplex DNA towards excellent catalysts.

Introduction

Enzymes, the catalysts of biological systems, can mediate a wide range of chemical reactions with high catalytic rates and perfect regio-/stereoselectivity.^{1,2} Cofactors usually play a crucial role in this process.¹ Metal complexes are one important type of cofactor that occurs in many natural proteins to mediate various reactions.³ In a biomimetic spirit, catalytic metal complexes have also been widely used as cofactors to combine covalently or non-covalently with biomacromolecules, such as proteins, apo-enzymes and antibodies, to construct artificial metalloenzymes.⁴⁻⁸ In particular, the chiral feature of biomacromolecules makes artificial metalloenzymes an ideal catalyst for enantioselective reactions.⁹⁻¹²

DNA has served as a new biological scaffold in constructing artificial metalloenzymes for enantioselective catalysis.¹²⁻¹⁴ Roefles and Feringa et al. found the cofactor Cu(II)-(4,4'-dimethyl-2,2'-dipyridyl) could be most compatible with double-stranded DNA (dsDNA).¹⁵ The resulting artificial dsDNA metalloenzyme proceeds with excellent enantioselectivity in a series of enantioselective reactions.¹⁶⁻²⁶ Recently, G-quadruplex DNA, due to its diverse and tunable chiral structures,²⁷⁻³¹ has become another candidate in constructing DNA metalloenzymes for enantioselective reactions.

Considerable efforts have been made to develop cofactors for G-quadruplex DNA, including a Cu(II)-bipyridine complex and its analogs by Moses et al.,³² Cu(II) ions by this group,³³⁻³⁶ and Cu(II) cationic porphyrin by Hennecke et al.³⁷ However, these G-quadruplex metalloenzymes only provide modest enantioselectivity for enantioselective reactions, which is most possibly due to the case that these cofactors cannot fit well with the G-quadruplex DNA binding pocket, and consequently the resulting catalytic centers are not able to provide suitable interactions with prochiral substrates to promote efficient enantioselectivity. Therefore, to develop a suitable cofactor with the best fit to the binding pocket provided by G-quadruplex DNA is a key issue in constructing efficient artificial G-quadruplex metalloenzymes.

Through the lens of molecular biology, G-quadruplex DNAs are attractive targets for the development of anticancer drugs.^{38,39} Small molecules capable of inducing and stabilizing G-quadruplex DNA structure in human telomeres and oncogene-promoter regions have shown therapeutic intervention in cancer.⁴⁰⁻⁴³ It is reported that a family of terpyridine-metal complexes shows good affinity and high selectivity for human telomeric G-quadruplex DNA.^{44,45} Inspired by this specific recognition, in this work we have employed terpyridine-Cu(II) complexes (CuLn) as cofactors to construct human telomeric G-quadruplex DNA metalloenzymes. Our study shows that the resulting human telomeric G-quadruplex DNA-CuL1 can afford excellent catalytic function in a Diels-Alder reaction with both high enantioselectivity over 99% enantiomeric excess (ee) and about 73-fold rate increase compared to CuL1 alone. To the best of our knowledge, this is the most efficient G-quadruplex DNA metalloenzyme for Diels-Alder reaction reported so far. Circular dichroism (CD) spectroscopic analysis, UV melting

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experiment and isothermal titration calorimetry (ITC) further indicate that the terpyridine-Cu(II) cofactor not only functions as a catalytic center but also plays an important role in inducing and stabilizing human telomeric G-quadruplex DNA with a catalytically active three-dimensional structure. Based on this proof-of-concept experiment, it is envisioned that much broader G-quadruplex metalloenzymes could be found by exploring G-quadruplex-targeting ligand pool.

Results

Terpyridine-Cu(II) complex targeting G-quadruplex DNA to produce artificial metalloenzyme

Inspired by the G-quadruplex DNA targeting nature of terpyridine-metal complexes, we synthesized the terpyridine-Cu(II) complex (CuL1) and employed CuL1 to directly interact with a human telomeric DNA sequence (HT21) (Fig. 1a). CD spectroscopic analysis show that, upon interaction of CuL1 with HT21, an obvious spectral change in the region of 220–310 nm is observed, accompanied by a new CD band ranging from 310–370 nm (Fig. 1b). The spectral region of 310–370 nm originates from the electric transition of the terpyridine-Cu(II) core.⁴⁶ Therefore, the new CD band (310–370 nm) can be readily ascribed to the induced circular dichroism (ICD) signal of CuL1 imposed by the chiral surroundings of HT21, which is also direct evidence of the CuL1-HT21 association. The CD spectrum of 220–310 nm is usually indicative of the G-quadruplex DNA structure.^{29–31} However, it is complicated in this system by the overlapping absorption band of CuL1. To aid in understanding the CD results, UV melting experiments were performed (Fig. 1c). The melting temperature (T_m) of HT21 increases by approximately 13 °C upon interaction with CuL1, which clearly indicates that the conformation of HT21 is significantly stabilized by the HT21-CuL1 complex formation. In this respect, the CD spectral change in the region of 220–310 nm contains much information on the structural transition of HT21. Moreover, the increased T_m clearly indicates that this structural transition occurs from a looser state to a more stable G-quadruplex DNA conformation. This deduction is further supported by the ITC experiment (Fig. 1d and 1e), which can determine the molecular nature of the binding interaction from a thermodynamic perspective.⁴⁷ The binding isotherms reveal a complex binding process and can be best fit by a three-event model (Fig. 1e and Table S1†).^{48,49} The high-affinity binding site allows the interaction of one CuL1 molecule with HT21 by an affinity constant of approximately 10^8 M^{-1} . Thermodynamic parameters further show that this strong association process is driven by a highly favorable enthalpy and a less unfavorable entropic contribution. The favorable enthalpy can be linked to either Hoogsteen hydrogen bond formation or association between CuL1 and HT21, which contributes to the structural regulation of HT21 from a looser state to a more stable G-quadruplex DNA conformation, whereas the unfavorable entropy change is mainly ascribed to decreased freedom from both the compact G-quadruplex DNA structure and the bound state of CuL1.⁵⁰

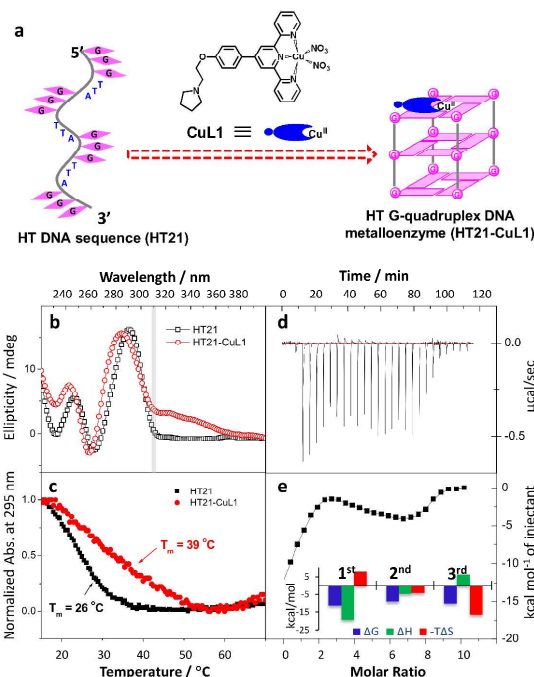



Fig. 1 Construction of human telomeric G-quadruplex DNA metalloenzyme. (a) Schematic diagram of molecular recognition between terpyridine-Cu(II) complex (CuL1) and human telomeric DNA sequence (HT21: 5'-(GGGTTA)₃GGG-3'). (b) CD spectra. (c) UV melting curves monitored by UV absorption at 295 nm. (d, e) Isothermal titration calorimetry (ITC) experiment. Raw ITC data is shown in panel (d); Plots of integrated calorimetric data after control subtraction (solid square), fitting curve (solid line) with a three-event binding model, and corresponding thermodynamic parameters are included in panel (e).

Upon confirming that CuL1 can directly induce and stabilize the HT21 G-quadruplex structure to form a stable HT21-CuL1 complex, we next select the Diels-Alder reaction between azachalcone (**1a**) and cyclopentadiene (**2**) as a model to test the catalytic performance of HT21-CuL1. As shown in Table 1, HT21-CuL1 exhibits much higher catalytic performance than CuL1 (entry 2 vs. entry 1), and catalyzes the reaction with 92% conversion (turnover number (TON) = 9.2), high diastereoselectivity of product **3a** (endo/exo of 97:3) and 90% ee of the major endo isomer in 24 hours (entry 2). Although these catalytic data were modest compared with the results with a dsDNA-based metalloenzyme,¹⁵ this result gives the best enantiomeric outcome concerning G-quadruplex DNA-based catalysis. This finding thus proves that CuL1 can act as an effective cofactor for the construction of a human telomeric G-quadruplex DNA metalloenzyme. More interestingly, combining the structural analysis and catalytic data, it is obvious that the CuL1 cofactor not only serves as a catalytic center but also shows structural regulation of human telomeric G-quadruplex DNA to promote excellent catalysis.

Structural optimization of G-quadruplex DNA in artificial metalloenzyme

In both natural and artificial metalloenzymes, subtle modifications of the secondary coordination sphere of the metal center usually exert a substantial influence on the



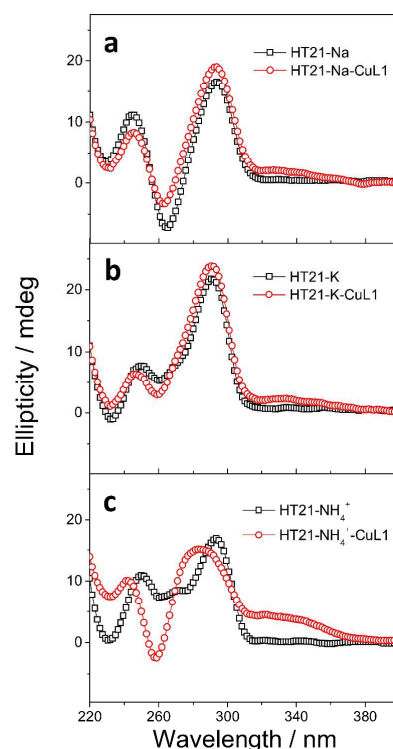
Table 1 Diels-Alder reaction catalyzed by human telomeric G-quadruplex DNA metalloenzyme


Entry	DNA Sequence (5'→3')	M ⁺	Conv. (%) ^a	endo/exo ^b	ee (% endo) ^b
1 ^c	-	-	42	71/29	0
2	HT21: (GGGTTA) ₃ GGG	-	92	97/3	90
<i>Conformation stabilized by monovalent cations</i>					
3	HT21:	Na ⁺	54	89/11	54
4	(GGGTTA) ₃ GGG	K ⁺	48	88/12	49
5		NH ₄ ⁺	99	99/1	94
<i>Flanking base variants</i>					
6	A-HT21: A(GGGTTA) ₃ GGG	NH ₄ ⁺	97	98/2	90
7	TA-HT21: TA(GGGTTA) ₃ GGG	NH ₄ ⁺	96	98/2	94
8	TTA-HT21: TTA(GGGTTA) ₃ GGG	NH ₄ ⁺	98	98/2	93
9	HT21-T: (GGGTTA) ₃ GGGT	NH ₄ ⁺	97	97/3	88
10	HT21-TT: (GGGTTA) ₃ GGGTT	NH ₄ ⁺	95	96/4	81
11	HT21-TTA: (GGGTTA) ₃ GGGTTA	NH ₄ ⁺	95	96/4	73

^a Determined for the crude product by HPLC analysis on a chiral stationary phase (Supplementary Methods Note 5), Reproducible within $\pm 2\%$. ^b Determined by chiral-phase HPLC. Reproducible within $\pm 2\%$. ^c Cu(L1)(NO₃)₂ alone as catalyst. Reaction conditions: **1a** (1 mM), **2** (10 μ L, 260 mM), human telomeric G-quadruplex DNA (50 μ M), Cu(L1)(NO₃)₂ (100 μ M), NaCl (50 mM) or KCl (150 mM) or NH₄Cl (30 mM), MOPS buffer (0.5 mL, 20 mM, pH 6.5), 4 $^{\circ}$ C, 24h.

catalytic function of the metalloenzyme.^{12,51} For the HT21-CuL1 metalloenzyme, this phenomenon provides an opportunity to improve the catalytic performance by varying the human telomeric G-quadruplex DNA structure. For this purpose, monovalent cations M⁺ (Na⁺/K⁺/NH₄⁺) were introduced to induce HT21 to form a G-quadruplex structure in advance,⁵² and then CuL1 was added to construct an artificial human telomeric G-quadruplex DNA metalloenzyme, represented by HT21-M⁺-CuL1.

The CD spectra of HT21-Na⁺-CuL1 (Fig. 2a) and HT21-K⁺-CuL1 (Fig. 2b) indicate that CuL1 has less structural regulation for HT21 G-quadruplex DNA (220–310 nm), and the ICD signal for CuL1 (310–370 nm) is also obscure. However, for HT21-NH₄⁺-CuL1, substantial structural change of HT21 G-quadruplex DNA (220–310 nm) and a clear ICD signal (310–370 nm) are observed (Fig. 2c). In addition, HT21-M⁺-CuL1 shows distinct catalytic behaviors for the Diels-Alder reaction compared with HT21-CuL1 (Table 1, entries 3–5 vs. entry 2). HT21-Na⁺-CuL1 shows an obvious decrease in both catalytic activity (conv. = 54%, TON = 5.4) and enantioselectivity (54% ee) (Table 1, entry 3). A similar case holds for HT21-K⁺-CuL1, with 48% conversion (TON = 4.8) and 49% ee (Table 1, entry 4). Surprisingly, when NH₄⁺

**Fig. 2** CD spectra of HT21 in the presence of different monovalent cations and their corresponding metalloenzyme. (a) 50 mM NaCl; (b) 150 mM KCl; (c) 30 mM NH₄Cl (DNA strand concentration, 5 μ M; CuL1, 10 μ M; MOPS buffer, 20 mM, pH 6.5).

was selected to stabilize HT21 G-quadruplex DNA, the HT21-NH₄⁺-CuL1 metalloenzyme demonstrated a substantial increase in catalytic activity compared with HT21-CuL1, showing 99% conversion (TON = 9.9) and excellent diastereoselectivity (endo/exo of 99:1), and the enantioselectivity for the major endo product **3a** increased to 94% ee (Table 1, entry 5). For a comparison of intrinsic catalytic activity, we performed an enzymatic dynamics study. The Michaelis-Menten kinetics showed that the HT21-NH₄⁺-CuL1 metalloenzyme reaches a catalytic efficiency (k_{cat}/K_M) of $36 \pm 4 \text{ M}^{-1}\text{s}^{-1}$ ($k_{\text{cat}} = 0.0015 \pm 0.0001 \text{ s}^{-1}$, $K_M = 41 \pm 3 \text{ }\mu\text{M}$), which is approximately 73-fold more active than CuL1 ($0.49 \pm 0.06 \text{ M}^{-1}\text{s}^{-1}$) (Fig. S1†). The rate acceleration in this system is comparable to dsDNA-based catalysis.⁵³ At the same time, HT21-NH₄⁺-CuL1 metalloenzyme shows 10-fold rate increase compared to HT21-CuL1 (Fig. S1†). Based on the above results, it is concluded that NH₄⁺ ion can function as an efficient chaperone for the HT21 G-quadruplex DNA metalloenzyme to create a favorable modification of HT21 G-quadruplex DNA structure, which can help the CuL1 cofactor to fit well with the binding pocket within HT21 G-quadruplex DNA and consequently promote the Diels-Alder reaction.

Under optimized experimental conditions in NH₄⁺ media, we further modified the HT21 G-quadruplex structure by introducing 1–3 flanking bases at the 5' or 3' end. These two positions were selected because the binding of terpyridine-Cu(II) complex with human telomeric G-quadruplex DNA



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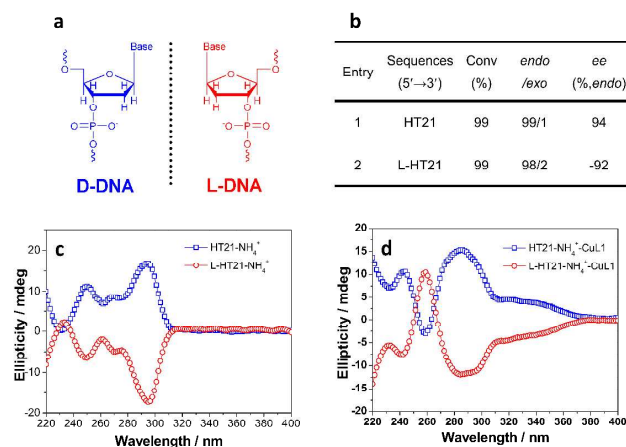


Fig. 3 Inversion of global chirality of G-quadruplex DNA metalloenzyme. (a) Natural D-DNA building block for HT21 and its enantiomer for L-HT21. (b) The Diels-Alder reaction (**1a** and **2**) catalyzed by HT21 and L-HT21 G-quadruplex DNA metalloenzymes. (c) CD spectra of HT21 and L-HT21 in NH_4^+ media (30 mM NH_4Cl). (d) CD spectra of HT21 G-quadruplex DNA metalloenzyme and L-HT21 G-quadruplex DNA metalloenzyme.

suggests that the aromatic surface around the copper could interact with human telomeric G-quadruplex DNA by π - π stacking on the external G-quartet.⁴⁴ Therefore, the flanking bases at the 5' or 3' end are also expected to modify the microenvironment of the CuL1 cofactor in the HT21-NH_4^+ - CuL1 metalloenzyme. For the Diels-Alder reaction between **1a** and **2**, these derivative human telomeric G-quadruplex DNA metalloenzymes present an interesting effect on the enantioselectivity (Table 1, entries 6-11). The sequences with flanking bases at the 5'-end have little effect on enantioselectivity (90-94% ee) compared with HT21-NH_4^+ - CuL1 (Table 1, entries 6-8 vs. entry 5). In contrast, a negative effect on enantioselectivity can be observed when the flanking bases are at the 3'-end (Table 1, entries 9-11 vs. entry 5). These data clearly indicate that the CuL1 cofactor is more sensitive to the 3'-end of HT21 G-quadruplex DNA, and a crowded microenvironment at the 3'-end is not favorable for high enantioselectivity.

The above data strongly suggest that the conformation of HT21 G-quadruplex DNA plays a dominant role in the enantioselective control. Therefore, based on the high level of enantioselectivity afforded by HT21-NH_4^+ - CuL1 , we further considered whether we could selectively access either enantiomer in the Diels-Alder reaction by tuning the HT21 G-quadruplex DNA conformation. To this end, mutation was performed by replacing the naturally occurring D-DNA with L-DNA (Fig. 3a). CD spectra show that L-HT21 forms a G-quadruplex structure that is a mirror image of natural HT21 (Fig. 3c). More importantly, upon assembly with CuL1 , the resulting L-HT21- NH_4^+ - CuL1 metalloenzyme also presents the mirror image structure of natural HT21-NH_4^+ - CuL1 (Fig. 3d). For the Diels-Alder reaction of **1a** and **2**, L-HT21- NH_4^+ - CuL1 can successfully switch the absolute configuration of the major product **3a** from 94% ee to -92% ee (Fig. 3b).

Structural optimization of cofactor structure in artificial metalloenzyme

Having validated the optimization of the second coordination sphere around Cu(II) by tuning the G-quadruplex structures, we further optimized the CuLn cofactor by tuning the terpyridine ligand, namely the first coordination sphere around Cu^{II} (Table 2). It is found that high catalytic activity and high enantioselectivity are not specific to CuL1 as a cofactor but are also obtained for other terpyridine- Cu(II) complexes possessing aminoalkyl side chains (CuLn , $n=2-4$) (Table 2, entries 1-4). In contrast, when the side chain (R) was changed from an aminoalkyl to a methyl group (CuL5), the HT21-NH_4^+ - CuL5 metalloenzyme showed substantial declines in both catalytic activity (conv. = 37%, TON = 3.7) and enantioselectivity (18% ee) (Table 2, entry 5). With further removal of the aromatic ring (CuL6), the HT21-NH_4^+ - CuL6 metalloenzyme led to only 25% conversion (TON = 2.5) and a complete inhibition of enantioselectivity (Table 2, entry 6). These catalytic data show that both the aminoalkyl side chain and the aromatic ring are key components for the terpyridine- Cu(II) cofactor to construct an efficient human telomeric G-quadruplex DNA metalloenzyme.

To clarify the underlying reason, we further focused on structural understanding of the HT21-NH_4^+ - CuLn metalloenzymes. We first used CD spectroscopy to detect the interaction of HT21-NH_4^+ with terpyridine- Cu(II) complexes possessing different side chains (Fig. 4). In comparison with HT21-NH_4^+ itself, all test terpyridine- Cu(II) cofactors demonstrate structural regulation on the HT21 G-quadruplex DNA structure (220-310 nm), but the degree of structural change is strongly dependent upon the type of side chain (R). At the same time, the ICD signals of the terpyridine- Cu(II) cofactor (310-370 nm) imposed by HT21 G-quadruplex DNA

Table 2 The ligand effect of terpyridine- Cu(II) cofactor on the catalytic function of human telomeric G-quadruplex DNA metalloenzyme

Entry	Ligands	Conv. (%) ^a	endo/exo ^b	ee (% endo) ^b
1	L1	99	99/1	94
2	L2	99	98/2	93
3	L3	99	97/3	92
4	L4	95	98/2	93
5	L5	37	81/19	18
6	L6	25	68/32	6

^a Determined for the crude product by HPLC analysis on a chiral stationary phase (Supplementary Methods Note 5). Reproducible within $\pm 2\%$. ^b Determined by chiral-phase HPLC. Reproducible within $\pm 2\%$. Reaction conditions: **1a** (1 mM), **2** (10 μL , 260 mM), HT21 (50 μM), $\text{Cu(Ln)(NO}_3)_2$ (100 μM), NH_4Cl (30 mM), MOPS buffer (0.5 mL, 20 mM, pH 6.5), 4 $^\circ\text{C}$, 24h.

Table 3 The binding thermodynamic parameters of different cofactors (CuLn) to human telomeric G-quadruplex DNA in 30 mM NH₄Cl at 298K

Cofactor	High-affinity binding site					Low-affinity binding site				
	K _{a1} (M ⁻¹)	n ₁	ΔH ₁ ^a	-TΔS ₁ ^a	ΔG ₁ ^a	K _{a2} (M ⁻¹)	n ₂	ΔH ₂ ^a	-TΔS ₂ ^a	ΔG ₂ ^a
CuL1 ^b	(1.7±0.2)E7	1.0	4.4±0.1	-14.3	-9.9	(6.6±0.5)E5	5.0	-2.2±0.1	-5.7	-7.9
CuL2 ^b	(1.7±0.3)E7	1.1	4.2±0.3	-14.0	-9.8	(1.0±0.1)E6	5.8	-2.1±0.1	-6.1	-8.2
CuL3 ^b	(1.7±0.3)E7	1.1	6.7±0.2	-16.6	-9.9	(1.1±0.1)E6	6.8	-1.5±0.1	-6.8	-8.3
CuL4 ^b	(1.7±0.3)E7	1.2	4.2±0.1	-14.1	-9.9	(5.8±0.6)E5	5.4	-2.5±0.1	-5.3	-7.8
CuL5 ^c	(2.0±0.5)E5	2.5	-1.1±0.1	-6.1	-7.2	-	-	-	-	-
CuL6 ^d	-	-	-	-	-	-	-	-	-	-

^a Units are kcal·mol⁻¹. ^b The data were obtained by the two-event binding model. ^c The data were obtained by the one-event binding model. ^d The affinity between the reactants is too low, neither the affinity nor the enthalpy of binding can be reliably determined.

also vary with different side chains. In detail, HT21-NH₄⁺-CuLn (n=1-4) metalloenzymes with terpyridine-Cu(II) cofactors possessing aminoalkyl side chains show similar CD spectral features, with clear structural regulation of the HT21 G-quadruplex DNA (220-310 nm) accompanied by the appearance of a strong ICD signal of the CuLn cofactors (Fig. 4a). Upon changing the cofactor to CuL5 (Fig. 4b), the structural transition of HT21 G-quadruplex DNA is also observed but shows a distinct CD feature from the HT21-NH₄⁺-CuLn (n=1-4) metalloenzyme. Moreover, HT21-NH₄⁺-CuL5 exhibits a smaller magnitude of ICD signal compared with HT21-NH₄⁺-CuLn (n=1-4). Upon changing to CuL6, the structural transition of HT21 becomes less notable, and the

ICD signal is also obscured (Fig. 4c). The ICD signal in the wavelength region 310-370 nm originates from the electric transition of the terpyridine-Cu(II) core, which is common to all six CuLn cofactors. Thus, the distinct amplitude of the ICD signal in HT21-NH₄⁺-CuLn indicates a different molecular orientation of the terpyridine-Cu(II) chromophore relative to the surrounding nucleobases,⁵⁴ which gives direct evidence of different recognition between CuLn and HT21-NH₄⁺ G-quadruplex DNA. More interestingly, a positive correlation between catalytic performance and amplitude of ICD signal indicates that the catalytic function of HT21-NH₄⁺-CuLn metalloenzyme is strongly dependent on the molecular recognition between CuLn and HT21 G-quadruplex DNA.

This conclusion based on CD data is further supported by ITC data (Table 3, Fig. S2† and Fig. S3†). For terpyridine-Cu(II) complexes possessing aminoalkyl side chains (CuLn, n=1-4), all binding isotherms are similar and reveal obvious two-event binding process. The high-affinity binding process, with a binding constant (K_{a1}) of 1.7×10⁷ M⁻¹, is the result of a highly favorable entropy change and a smaller unfavorable enthalpy change. The low-affinity binding process displays a binding constant (K_{a2}) of 5.8×10⁵ ~1.1×10⁶ M⁻¹, which is 16~30 times lower than the K_{a1}. In contrast to the high-affinity binding process, this low-affinity binding is favored by both entropy and enthalpy changes. The titration of CuL5 into HT21-NH₄⁺ displays a relatively simple binding isotherm. A binding constant (K_a) of 2.0×10⁵ M⁻¹ is estimated by fitting a single binding event model, which is of the same order of magnitude as the low-affinity binding of CuLn (n=1-4) to HT21-NH₄⁺ G-quadruplex DNA. Furthermore, the molecular recognition process between CuL5 and HT21-NH₄⁺ G-quadruplex DNA also shows a similar thermodynamic signature to the low-affinity binding of CuLn (n=1-4) to HT21-NH₄⁺ G-quadruplex DNA (Fig. S3†). In contrast, the binding isotherm of CuL6 to HT21-NH₄⁺ is essentially the same as the background calorimetric titration of MOPS buffer to HT21-NH₄⁺, suggesting that weak binding between CuL6 and HT21-NH₄⁺ occurs, and neither the binding constant nor the thermodynamic parameters can be reliably determined. Thus, based on the different energetic characteristics, the ITC results clearly show three types of molecular recognition processes, and correspondingly, these three distinct interaction processes give the binding affinity order of CuLn (n=1-4)>CuL5>CuL6 to HT21-NH₄⁺ G-quadruplex

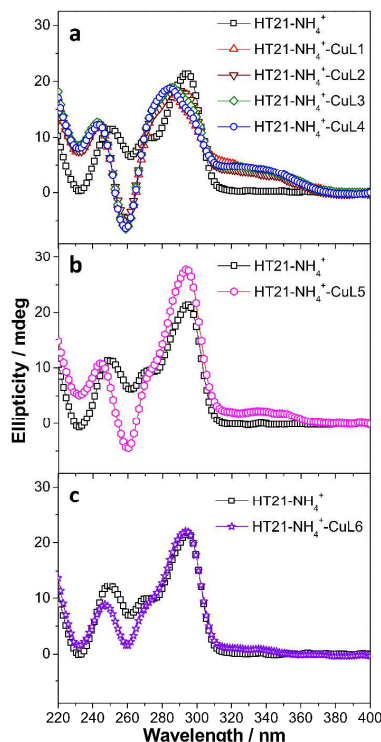


Fig. 4 CD spectra of human telomeric G-quadruplex DNA metalloenzymes with different terpyridine-Cu(II) cofactors compared with HT21 itself in NH₄⁺ media. (a) CuLn (n=1-4); (b) CuL5; (c) CuL6 (DNA strand concentration, 5 μM; NH₄Cl, 30 mM; CuLn, 10 μM; MOPS buffer, 20 mM, pH 6.5).



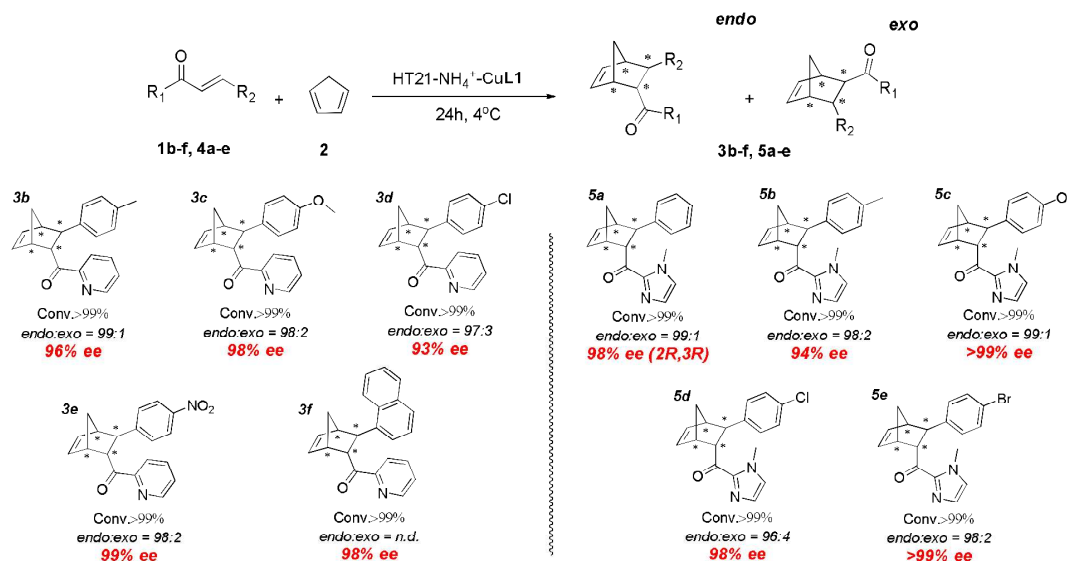


Fig. 5 Substrate scope for human telomeric G-quadruplex DNA metalloenzyme. Reaction conditions: 1 or 4 (1 mM), 2 (10 μ L), HT21 (50 μ M), CuL1 (100 μ M), NH_4Cl (30 mM), MOPS buffer (20 mM, pH 6.5), 4 $^\circ C$, 24h. All data are averaged over two experiments and reproducible within $\pm 2\%$. Conversion measured by 1H -NMR analysis of the crude product. The endo/exo ratios and ee (for the endo isomer) are determined by chiral-phase HPLC. The absolute configuration of 5a was obtained by referring 16.

DNA. This sequence is consistent with not only the amplitude of the ICD signal in the CD experiment discussed above but also the ΔT_m values in UV melting experiments, where $CuLn$ ($n=1-4$) increases the melting temperature of $HT21-NH_4^+$ most significantly ($\Delta T_m \approx 20$ $^\circ C$), followed by $CuL5$ ($\Delta T_m \approx 10$ $^\circ C$) and $CuL6$ ($\Delta T_m \approx 5$ $^\circ C$) (Fig. S4 †).

In connection with the catalytic data and the above characterizations (CD, UV melting, and ITC), we thus concluded that the molecular recognition between HT21 G-quadruplex DNA and terpyridine- $Cu(II)$ plays an essential role in the catalytic function of the $HT21-NH_4^+-CuLn$ metalloenzyme. In fact, the high affinity of $CuLn$ ($n=1-4$) for HT21 G-quadruplex DNA can be easily understood as the interplay of two effects: 1) the $\pi-\pi$ stacking interactions between the terpyridine and the G-quartet; 2) the electrostatic interaction between the positive charge of the aminoalkyl group and the negative charge of the phosphate backbones.^{44,55}

Substrate Scope

With the excellent $HT21-NH_4^+-CuL1$ metalloenzyme in hand, we further tested the substrate specificity (Fig. 5). The $HT21-NH_4^+-CuL1$ metalloenzyme was found to be an active catalyst for all the tested aza-chalcone substrates (1b-f). In general, the introduction of a bulkier R_2 group has a positive influence on the enantioselectivity. In particular, R_2 with a nitro group afforded up to 99% ee (3e). Furthermore, high activity and enantioselectivity are also observed for the other type of dienophiles possessing acyl-imidazol groups (4a-e). A nearly pure isomers (> 99% ee) are obtained for 4-MeO and 4-Br (5c and 5e).

Discussion

Small molecule targeting of G-quadruplex DNA represents an attractive approach to anticancer drug design.⁴⁰⁻⁴³ This recognition process is analogous to the assembly of an artificial enzyme between a catalytic cofactor and a biomolecular scaffold.¹⁰⁻¹² Following this idea, we have found an excellent G-quadruplex metalloenzyme through molecular recognition between terpyridine- $Cu(II)$ and human telomeric G-quadruplex DNA, although the precise nature of the interaction is not well known. This modular assembly method allows rationally tunable enantioselectivity of the Diels-Alder reaction. The G-quadruplex DNA scaffold can be modulated by the addition of cation, modification of flanking bases at the 3'-end, and even inversion of global chirality by replacing natural *D*-DNA with unnatural *L*-DNA. Modification of the substitute in the terpyridine ligand can also change the binding affinity between G-quadruplex DNA and the terpyridine- $Cu(II)$ cofactor, which alters the microenvironment of the active site. Based on this proof-of-concept experiment, we expect this approach to provide a general way to find a broad variety of G-quadruplex metalloenzymes by screening G-quadruplex-targeting ligand pools.

Conclusions

In conclusion, we have designed a G-quadruplex metalloenzyme that can afford both high catalytic activity and high enantioselectivity (up to 99% ee) for the Diels-Alder reaction. The success of this artificial metalloenzyme can be ascribed to the G-quadruplex DNA-targeting nature of terpyridine- $Cu(II)$. The terpyridine- $Cu(II)$ cofactor is found to play dual roles as both an active site for chemical catalysis and a structural regulator for the folding of the catalytic DNA structure. A positive correlation between the affinity of



terpyridine-Cu(II) for HT21 and the enantioselectivity of the Diels-Alder reaction is observed, which further confirms the importance of molecular recognition in constructing artificial G-quadruplex DNA metalloenzymes. Our work paves a new way to design G-quadruplex-based DNAzymes by learning from G-quadruplex-targeting drug design.

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Notes and references

- 1 T. Bugg, *Introduction to enzyme and coenzyme chemistry*, Blackwell Pub., Oxford, UK ; Malden, MA, USA, 3rd edn., 2012.
- 2 W. D. Fessner and T. Anthonsen, *Modern biocatalysis : stereoselective and environmentally friendly reactions*, Wiley-VCH, Weinheim, 2009.
- 3 D. E. Metzler and C. M. Metzler, in *Biochemistry : the chemical reactions of living cells*, Harcourt/Academic Press, San Diego, Calif., 2nd edn., 2001, vol. 1, ch. 16, pp. 837.
- 4 K. L. Harris, S. Lim and S. J. Franklin, *Inorg. Chem.*, 2006, **45**, 10002.
- 5 Y. Lu, N. Yeung, N. Sieracki and N. M. Marshall, *Nature*, 2009, **460**, 855.
- 6 M. Dürrenberger and T. R. Ward, *Curr. Opin. Chem. Biol.*, 2014, **19**, 99.
- 7 I. D. Petrik, J. Liu and Y. Lu, *Curr. Opin. Chem. Biol.*, 2014, **19**, 67.
- 8 F. Yu, V. M. Cangelosi, M. L. Zastrow, M. Tegoni, J. S. Plegaria, A. G. Tebo, C. S. Mocny, L. Ruckthong, H. Qayyum and V. L. Pecoraro, *Chem. Rev.*, 2014, **114**, 3495.
- 9 M. T. Reetz, *Angew. Chem. Int. Ed.*, 2011, **50**, 138.
- 10 R. Krämer, *Angew. Chem. Int. Ed.*, 2006, **45**, 858.
- 11 T. R. Ward, *Acc. Chem. Res.*, 2011, **44**, 47.
- 12 J. Bos and G. Roelfes, *Curr. Opin. Chem. Biol.*, 2014, **19**, 135.
- 13 A. Jäschke, in *Molecular encapsulation: organic reactions in constrained systems*, John Wiley & Sons, Ltd, 2010, ch. 14, pp. 377.
- 14 S. Park and H. Sugiyama, *Angew. Chem. Int. Ed.*, 2010, **49**, 3870.
- 15 G. Roelfes, A. J. Boersma and B. L. Feringa, *Chem. Commun.*, 2006, 635.
- 16 A. J. Boersma, B. L. Feringa and G. Roelfes, *Org. Lett.*, 2007, **9**, 3647.
- 17 D. Coquière, B. L. Feringa and G. Roelfes, *Angew. Chem. Int. Ed.*, 2007, **46**, 9308.
- 18 E. W. Dijk, B. L. Feringa and G. Roelfes, *Tetrahedron: Asymmetry*, 2008, **19**, 2374.
- 19 A. J. Boersma, B. L. Feringa and G. Roelfes, *Angew. Chem. Int. Ed.*, 2009, **48**, 3346.
- 20 N. Shibata, H. Yasui, S. Nakamura and T. Toru, *Synlett*, 2007, 1153.
- 21 Y. H. Li, C. H. Wang, G. Q. Jia, S. M. Lu and C. Li, *Tetrahedron*, 2013, **69**, 6585.
- 22 J. Wang, E. Benedetti, L. Bethge, S. Vonhoff, S. Klusmann, J.-J. Vasseur, J. Cossy, M. Smietana and S. Arseniyadis, *Angew. Chem. Int. Ed.*, 2013, **52**, 11546.
- 23 L. Gjonaj and G. Roelfes, *ChemCatChem*, 2013, **5**, 1718.
- 24 H. Zhao and K. Shen, *RSC Adv.*, 2014, **4**, 54051.
- 25 E. Benedetti, N. Duchemin, L. Bethge, S. Vonhoff, S. Klusmann, J.-J. Vasseur, J. Cossy, M. Smietana and S. Arseniyadis, *Chem. Commun.*, 2015, **51**, 6076.
- 26 A. Draksharapu, A. J. Boersma, M. Leising, A. Meetsma, W. R. Browne and G. Roelfes, *Dalton Trans.*, 2015, **44**, 3647.
- 27 S. Burge, G. N. Parkinson, P. Hazel, A. K. Todd and S. Neidle, *Nucleic Acids Res.*, 2006, **34**, 5402.
- 28 J. Dai, M. Carver and D. Yang, *Biochimie*, 2008, **90**, 1172.
- 29 J. Kypr, I. Kejnovská, D. Renčíuk and M. Vorlíčková, *Nucleic Acids Res.*, 2009, **37**, 1713.
- 30 D. M. Gray, J. D. Wen, C. W. Gray, R. Repges, C. Repges, G. Raabe and J. Fleischhauer, *Chirality*, 2008, **20**, 431.
- 31 S. Masiero, R. Trotta, S. Pieraccini, S. De Tito, R. Perone, A. Randazzo and G. P. Spada, *Org. Biomol. Chem.*, 2010, **8**, 2683.
- 32 S. Roe, D. J. Ritson, T. Garner, M. Searle and J. E. Moses, *Chem. Commun.*, 2010, **46**, 4309.
- 33 C. H. Wang, G. Q. Jia, J. Zhou, Y. H. Li, Y. Liu, S. M. Lu and C. Li, *Angew. Chem. Int. Ed.*, 2012, **51**, 9352.
- 34 C. H. Wang, Y. H. Li, G. Q. Jia, Y. Liu, S. M. Lu and C. Li, *Chem. Commun.*, 2012, **48**, 6232.
- 35 C. H. Wang, G. Q. Jia, Y. H. Li, S. F. Zhang and C. Li, *Chem. Commun.*, 2013, **49**, 11161.
- 36 Y. H. Li, G. Q. Jia, C. H. Wang, M. P. Cheng and C. Li, *ChemBioChem*, 2015, **16**, 618.
- 37 M. Wilking and U. Hennecke, *Org. Biomol. Chem.*, 2013, **11**, 6940.
- 38 H. Y. Han and L. H. Hurley, *Trends Pharmacol. Sci.*, 2000, **21**, 136.
- 39 S. Neidle, *FEBS J.*, 2010, **277**, 1118.
- 40 D. J. Patel, A. T. Phan and V. Kuryavyy, *Nucleic Acids Res.*, 2007, **35**, 7429.
- 41 A. Arola and R. Vilar, *Curr. Top. Med. Chem.*, 2008, **8**, 1405.
- 42 D. Z. Yang and K. Okamoto, *Future Med. Chem.*, 2010, **2**, 619.
- 43 N. W. Luedtke, *Chimia*, 2009, **63**, 134.
- 44 H. Bertrand, D. Monchaud, A. De Cian, R. Guillot, J.-L. Mergny and M.-P. Teulade-Fichou, *Org. Biomol. Chem.*, 2007, **5**, 2555.
- 45 S. N. Georgiades, N. H. Abd Karim, K. Suntharalingam and R. Vilar, *Angew. Chem. Int. Ed.*, 2010, **49**, 4020.
- 46 I. Eryazici, C. N. Moorefield and G. R. Newkome, *Chem. Rev.*, 2008, **108**, 1834.
- 47 I. Haq, B. Z. Chowdhry and T. C. Jenkins, *Methods Enzymol.*, 2001, **340**, 109.
- 48 V. H. Le, R. Buscaglia, J. B. Chaires and E. A. Lewis, *Anal. Biochem.*, 2013, **434**, 233.
- 49 J. S. Hudson, L. Ding, V. Le, E. Lewis and D. Graves, *Biochemistry*, 2014, **53**, 3347.
- 50 D. H. Williams, E. Stephens, D. P. O'Brien and M. Zhou, *Angew. Chem. Int. Ed.*, 2004, **43**, 6596.
- 51 A. J. Kirby, *Angew. Chem. Int. Ed. Engl.*, 1996, **35**, 707.
- 52 N. V. Hud and J. Plavec, in *Quadruplex Nucleic Acids*, eds. S. Neidle and S. Balasubramanian, RSC, 2006, pp. 100.
- 53 A. J. Boersma, J. E. Klijn, B. L. Feringa and G. Roelfes, *J. Am. Chem. Soc.*, 2008, **130**, 11783.
- 54 M. Eriksson and B. Norden, *Methods Enzymol.*, 2001, **340**, 68.
- 55 C. Wei, L. Ren and N. Gao, *Int. J. Biol. Macromol.*, 2013, **57**, 1.

