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An oxidatively damaged G-quadruplex/hemin DNAzyme

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Abstract: Oxidative damage of guanine to 8-oxoguanine triggers a partial and variable loss of G-quadruplex/hemin DNAzyme activity and provide clues about the mechanistic origins of DNAzyme deactivation, which originates in an interplay between decreased G-quadruplex stability, lower hemin affinity and a modification of the nature of hemin binding site.

G-quadruplexes (G4s), assembled by four Gs in the same plane and more quarters subsequently self-stack *via* π -stacking interactions (Fig. 1), have recently attracted great interest for their possible biological roles, given that they are widely distributed in the human genomes and transcriptomes¹⁻⁴. G4s are also key players in biotechnology: thanks to their controllable stability and polymorphism⁵. G4s have been instrumental in the development of a new class of functional nucleic acids, chiefly the G4-DNAzymes (Fig. 2)⁵⁻⁸. The exquisite interaction between G4s and hemin (iron (III)-protoporphyrin IX) results in peroxidase-mimicking G4-DNAzyme systems⁹, which have been used as biosensors^{10,11} and biodevices^{5,12,13}. Over the past years, various strategies have been applied to enhance the enzymatic proficiency of G4-DNAzymes, including the addition of a proximal adenine or cytosine nucleotide to the 3' G-quartet of a parallel G4 (the preferential hemin binding site)¹⁴⁻²¹, the covalent linking of a substrate aptamer to G4 sequences²²⁻²⁴, or the addition of small-molecular catalytic enhancers²⁵⁻³⁰.

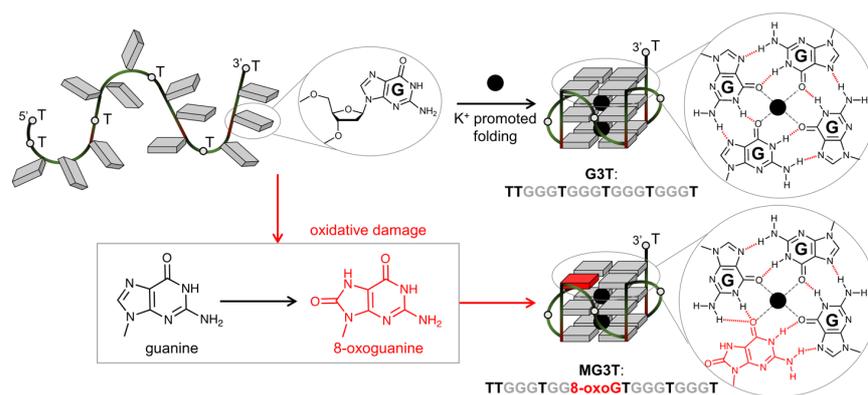


Fig. 1 Schematic representation of a parallel quadruplex-forming sequence (here, $T_2G_3TG_3TG_3TG_3T$) under its unfolded (left) and K^+ -promoted folded structure (i.e., a G-quadruplex, or G4, right), with detailed chemical structure of a guanine (G) and a G-quartet (right). Lower panel: oxidative damage of a guanine (G), converted into an 8-oxoguanine (8-oxoG), and consequences for the formation of an 8-oxoG-containing G-quartet.

Interestingly, most if not all G4-DNAzymes described to date are fueled by hydrogen peroxide (H_2O_2) as the stoichiometric oxidant⁹. In living organisms, H_2O_2 is known to oxidize DNA, notably converting Gs into 8-oxoguanine (8-oxoG, Fig. 1)³¹⁻³⁴. It was thus of interest to study the possible side-effect of H_2O_2 on the structural integrity of the G4 pre-catalyst. First insights have been provided by Sen *et al.* who showed by a combination of chemical probing and electron paramagnetic resonance (EPR) that G4s are indeed damaged by H_2O_2 , likely to be responsible for the inactivation of DNAzyme activity³⁵. More recently, Podbevsek *et al.* reported that a G4 structure is stable enough to tolerate a substitution of a guanine by an 8-oxoG³⁶.

We thus decided to systematically investigate the impact of oxidative lesions on enzymatic activity of parallel G4/hemin DNAzymes (Fig. 2). To this end, we designed a series of eleven G4-forming sequences without (controls) and with 8-oxoGs at various positions within the sequences belonging to the 3'-end quartet, the preferred hemin binding pocket (Fig. 2, Table 1); G4s with 8-oxoG located on the 5'-end and in the middle quartets were used as controls, as shown below. Three unmodified sequences were used as controls: G3T and the two related G3TC and G3A, in which a d(C) was added at the 3'-end (G3TC) or in which the final d(T) was replaced by a d(A) (G3A). G3T was selected as a model G4 given that its structure has been confirmed by nuclear magnetic resonance (NMR) spectroscopy (parallel G4, PDB ID: 2LK7)³⁷. 8-oxoG were introduced at positions 5, 9, 13 and 17 of both G3TC and G3A, thus providing a series of eight oxidatively modified sequences referred to as TC-5, TC-9, TC-13, TC-17, A-5, A-9, A-13, A-17, respectively (Fig. 2, Table 1).

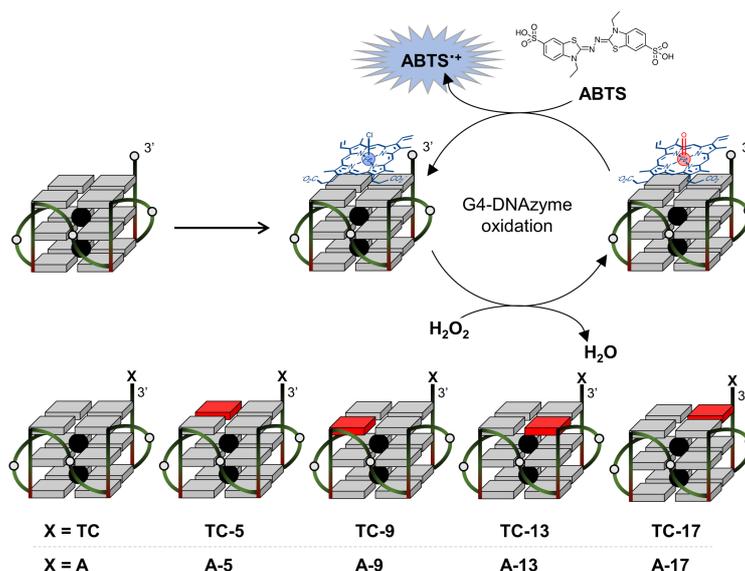


Fig. 2 Schematic representation of a G4/hemin DNAzyme system (upper panel) that triggers H_2O_2 -promoted oxidation of ABTS, and of 8-oxoG-containing G4s investigated in this study (8-oxoG in red, lower panel).

To demonstrate that all sequences adopt a G4 structure, thermal difference spectra (TDS)³⁸ were recorded. As seen in Fig. 3a, all folded sequences exhibited positive and negative peaks at 279 and 295 nm, respectively, in agreement with G4 formation, with or without oxidative damage. These results were consistent with previous reports that G4s can withstand G-to-8-oxoG mutations^{36,39,40}. Next, circular dichroism (CD) spectra showed that the incorporation of 8-oxoG in G3TC and G3A preserved the parallel topology of the parent sequences, as evidenced by positive and negative peaks at *ca.* 265 and *ca.* 242 nm, respectively (Fig. 3b)³⁹⁻⁴¹. The thermal stabilities of the resulting G4s were determined by UV-melting/annealing experiments (Fig. S1, Supporting Information): the melting temperature values (T_m , summarized in Table 1) were in line with previous reports^{36,39,40} that the presence of an 8-oxoG within a G4-forming sequence decreased the overall stability of the resulting G4 structure. Our results also highlight that G4s with a single 8-oxoG are quite stable, with T_m values >52 °C. Additionally, the four G3TC-related G4s have similar stabilities, as well as the four G3A-related G4s, which is suggestive of a similar hydrogen bonding network in these structures. Collectively, these results show that the 8-oxoG-modified G4s studied here tolerate oxidative damage of the guanine residues, both in terms of topology (a parallel fold is preserved) and stability (T_m decreased moderately and still above physiological temperature). As demonstrated by previously reported high-resolution structures (PDB IDs: 6IA0, 6IA4)³⁶, this stability could originate in the involvement of the oxygen of $\text{C}_6=\text{O}_6$ from the 8-oxoG in two H-bonds with the proximal G (Fig. 1), thereby participating to the stability of the 8-oxoG•G•G•G quartet, yet less stable than the native G•G•G•G quartet.

Table 1 DNA sequences information^a

Name	Sequences (5'→3')	T_m (°C) ^b	V_o (nM/s) ^c
Unmodified			
	5 9 13 17 ^d		
G3T	TT GGG T GGG T GGG T GGG T	>85	28.8±1.9
G3TC	TT GGG T GGG T GGG T GGG TC	>85	276.7±22.7
G3A	TT GGG T GGG T GGG T GGG A	>85	125.5±17.0
Modified by 8-oxoG			
TC-5	TT GGX T GGG T GGG T GGG TC	55.1	120.9±3.6
TC-9	TT GGG T GGX T GGG T GGG TC	52.4	189.5±21.3
TC-13	TT GGG T GGG T GGX T GGG TC	54.1	221.6±21.6
TC-17	TT GGG T GGG T GGG T GGX TC	54.2	57.6±4.6
A-5	TT GGX T GGG T GGG T GGG A	58.0	59.6±4.7
A-9	TT GGG T GGX T GGG T GGG A	57.7	87.8±11.3
A-13	TT GGG T GGG T GGX T GGG A	57.3	99.4±13.1
A-17	TT GGG T GGG T GGG T GGX A	58.0	73.3±4.5

^a X indicates that guanine is replaced by 8-oxoG. ^b T_m is obtained from the UV-melting experiments (Fig. S1). ^c Initial reaction velocity (V_o) is obtained from the DNAzyme activity assay (Fig. S2). ^d Nucleotide position.

In our modified sequences, G-to-8-oxoG substitutions are located at the 3' quartet, *i.e.*, close to the hemin binding site. We thus decided to assess whether and how the presence of 8-oxoG affects the catalytic proficiency of the resulting G4s. To this end, we implemented the well-studied oxidation of 2,2'-azino-bis(3-ethylbenzothiazoline-6-sulphonic acid) (ABTS) by H₂O₂ as a model reaction. This reaction provides an easily readable output, the evolution of absorbance of the oxidized product ABTS⁺ at $\lambda = 420$ nm as a function of time (Fig. S2). Several conclusions can be drawn from the initial reaction velocities (V_o), presented in Fig. 4a: *i-* the addition of d(TC) or d(A) at 3' end of a G4 improves its activity ($V_o = 29, 277$ and 126 nM/s for G3T, G3TC and G3A, respectively), which is consistent with previous reports¹⁵⁻¹⁹; *ii-* the presence of 8-oxoG decreases the catalytic efficiency (*e.g.*, $V_o = 121$ and 59 nM/s for TC-5 and A-5, respectively); but *iii-* the activity of 8-oxoG-modified G4s is still noticeable, being higher than that of hemin alone (blank, $V_o = 1.1$ nM/s) and of the parent G3T ($V_o = 29$ nM/s). The position of the 8-oxoG modification influences the catalysis (Fig. 4b), with lower catalytic efficiency for G4 bearing modifications close to 3' end of d(TC) and d(A) tails, ($V_o = 121, 58, 60$ and 73 nM/s for TC-5, TC-17, A-5 and A-17, respectively) as compared to G4s with modifications away from the tails ($V_o = 190, 223, 88$ and 99 nM/s for TC-9, TC-13, A-9 and A-13, respectively). To gain further insights

into the possible roles of oxidized Gs on catalysis, 8-oxoG were introduced both on the 5'-end (TC-15, Fig. S3) and in the middle G-quartet (TC-16, Fig. S3), which are not supposed to interact directly with hemin. Both sequences form G4 structures, as evidenced by TDS and CD, but TC-15 was found far more stable than TC-16 ($T_m = 55$ and 42°C , respectively, Fig. S3). The performance of TC-15 was comparable to that of the parent G3TC ($V_o = 259$ and 277 nM/s, respectively), in line with the preferential hemin binding site located on the 3'-end quartet¹⁴⁻²¹. The catalytic proficiency of TC-16 was weak ($V_o = 117$ nM/s), which may originate from its low thermal stability.

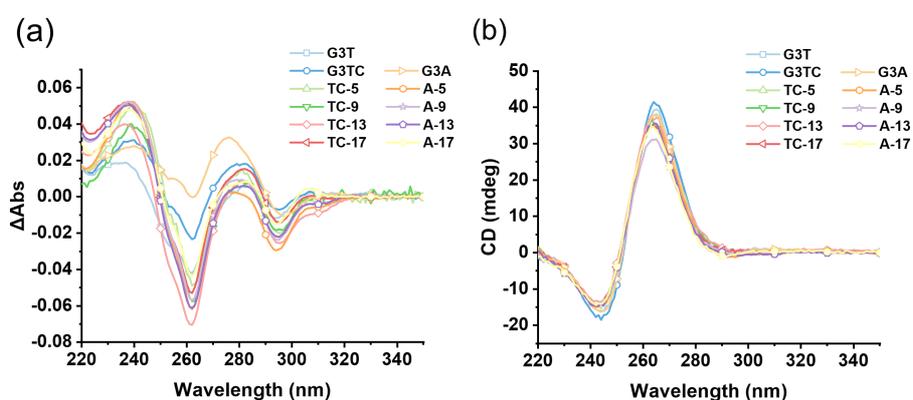


Fig. 3 Evidence for G-quadruplex formation. (a) Thermal difference spectra (TDS) of all sequences. Unless otherwise stated, UV-visible absorbance spectra were carried out in 10 mM pH 7.0 Tris-HCl containing 50 mM KCl and 5.0 μM DNA. TDS were the differences of absorbance between the high and low temperatures. Potassium concentration in solution was decreased to 5.0 mM because of the extremely high stabilities of G3T, G3A and G3TC. (b) CD spectra of all sequences in the presence of 50 mM KCl.

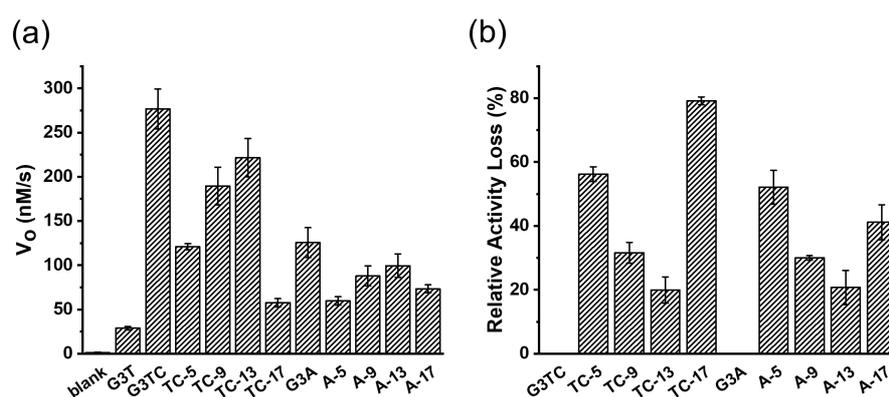


Fig. 4 Catalytic activities of DNAzyme without or with an oxidative lesion. (a) Initial reaction velocities (V_o , nM/s). Raw data are presented in Fig. S2. 'Blank' represents the V_o of hemin alone. Reaction conditions: 0.8 μM DNA, 0.4 μM hemin, 0.05 % Triton X-100, 2.0 mM ABTS and 0.6 mM H_2O_2 , 10 mM pH 7.0 Tris-HCl containing 50 mM KCl, 25°C . Error bars represented the standard deviation in three independent experiments. (b) Relative activity (V_o) loss after modification of G-quadruplexes, compared to the parent G3TC or G3A sequence, with no oxidation product.

Mechanistically, the activity of G4-DNAzyme complexes relies on the transient formation of high valent iron after the addition of H_2O_2 (Fig. 5a). These unstable intermediates can either oxidize ABTS (to form ABTS^+) or degrade the G4/hemin complex, or both. This was investigated by UV-Vis spectroscopy: a sharp Soret band (absorbance at $\lambda = 404$ nm) belonging to the hemin spectra appears upon addition of G4s (Fig. S4), indicating that both unmodified and modified G4s can coordinate the ferric ion. The degradation of G4/hemin complex by H_2O_2 was monitored *via* the time-dependent evolution of this Soret band in the absence of ABTS (Fig. 5a and S5): the good correlation between the apparent initial degradation rates (V_d , Fig. 5b) and V_o suggests that both unmodified and modified G4s activate hemin *via* a similar coordination pattern. These results showed again that modifications close to the sequence tails impair the catalytic activity of the resulting G4s more than modifications in the central positions (*e.g.*, $V_d = 4.6, 3.1$ and $6.9, 7.2$ nM/s for TC-5, TC-17 and TC-9, TC-13, respectively). Finally, the binding affinity (K_a) of hemin for G4s determined *via* UV-Vis titration (Table S1, Fig. 5b) demonstrated that 8-oxoG modifications decrease binding affinity, which may also contribute to the global loss of DNAzyme activity. Our results are in line with previous reports notably from Sen *et al.* who detected an oxidatively damaged G in a G4/hemin complex upon addition of H_2O_2 ³⁶, and Shangguan *et al.* who demonstrated that the inactivation of G4/hemin complex can be mainly attributed to the hemin degradation by H_2O_2 ⁴². We confirm that oxidative damage of G4 results in partial G4-DNAzyme inactivation (20-80%, Fig. 4b) and that both H_2O_2 -mediated degradation of hemin and oxidation of G4s contribute to the overall inactivation of the G4-DNAzyme system (Fig. 5a).

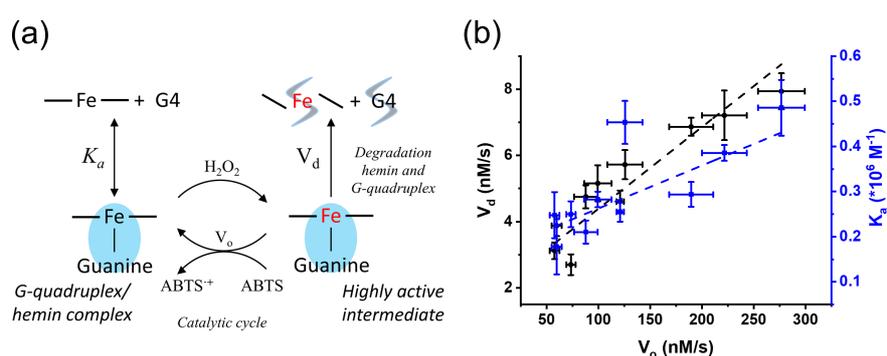


Fig. 5 (a) Scheme of the interaction between G-quadruplex and hemin (K_a), catalytic cycle (V_o) and degradation of G-quadruplex/hemin complex (V_d) in the absence of reductant substrate. Inactivation results from the degradation of both hemin and G-quadruplex. (b) Apparent initial reaction velocity (V_o) as functions of initial degradation velocity (V_d , black dot and left scale) and apparent binding constant between G-quadruplex and hemin (K_a , blue square and right scale). V_o was extracted from the above mentioned activity assay (Fig. S2). V_d is the initial degradation rate of G-quadruplex/hemin complex by H_2O_2 in the absence of reductant substrate (Fig. S5).

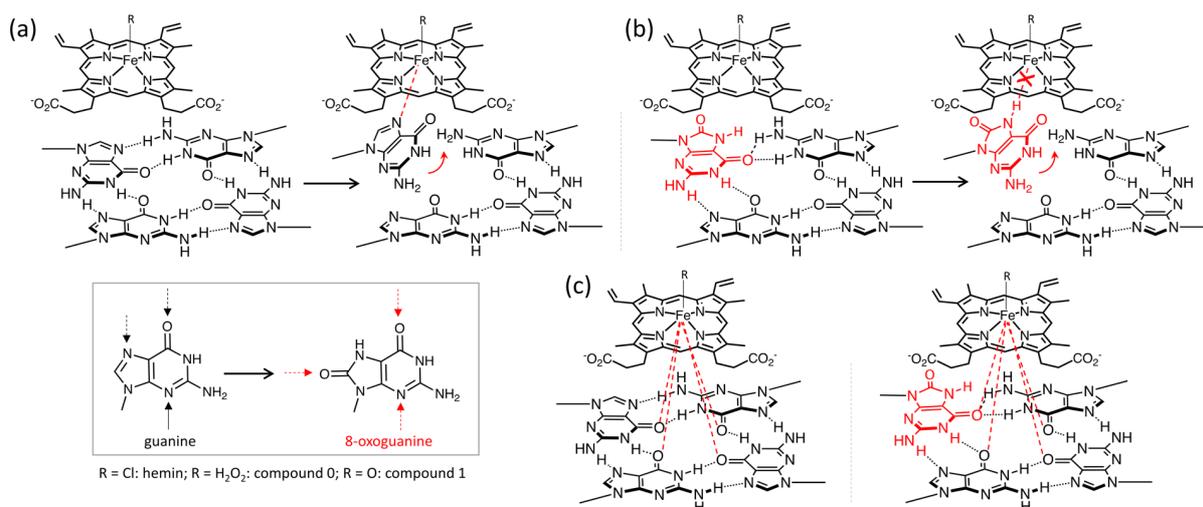


Fig. 6 Postulated intermediate for G4-DNAzyme catalysis: the guanine flipping model (a,b) versus the pan-coordination model (c). Inset: schematic representation of the coordination possibilities of G and 8-oxoG (arrows).

Finally, we analysed the coordination properties of both G and 8-oxoG. As seen in Fig. 6, while both of them have three possible coordination sites (dashed arrows), the most notable difference concerns the N₇-coordination, which is blocked in 8-oxoGs. This may have important consequence for coordinating hemin during the catalysis. Indeed, two catalytic intermediate models are currently postulated (Fig. 6a-c): in the first model, a G can transiently flip out from the G-quartet plane^{19,26,35,43-45} to coordinate the iron atom and activate it; in the second model, the iron atom coordination occurs *via* the oxygen atoms of the four carbonyl groups without G-quartet disruption. Our results suggest the reliability of the first model (Fig. 6b): even if the 8-oxoG can flip out of the modified G-quartet, it will not coordinate properly the iron atom given that its N₇ position three G can still coordinate to iron for the formation of activity intermediates. For the second model, the coordination between oxygen and iron seems to be not affected the modification of G-quartet (Fig. 6c), which is not compatible with activity loss herein. Of note, another possible explanation could involve a strand-slipped intermediate,⁴⁶ but no information has yet been collected to support it during DNAzyme-type catalysis.

Collectively, the sequential G-to-8-oxoG modification strategy described here confirms that DNA oxidation globally alters the catalytic proficiency of G4-based DNAzyme. They also provide invaluable insights into the mechanistic origins of activity loss, which seems to result from an interplay between decreased G4 stability, lower hemin affinity and a modification of the nature of the hemin binding pocket. Our results favour a guanine-flipping model, which is close to

what has been recently demonstrated as a naturally-occurring intermediate of G4-resolving helicase. This demonstration has profound implications for the design of ever-more efficient and stable G4-DNAzyme systems, which should now embed structurally flexible external quartet to allow for an optimized activation of the hemin iron atom while maintaining good π -stacking interactions with the macrocyclic scaffold of the hemin. This represents a novel example of a finely tuned and highly sophisticated mechanism that nature was able to create.

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Conflicts of interest

The authors declare no conflict of interest.

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