We demonstrate here that 8-amino guanine (R) strongly accelerates quadruplex formation, which makes this nucleobase the most attractive replacement for guanine in the context of tetramolecular parallel quadruplexes.

G-Quadruplexes are a family of four-stranded DNA structures stabilized by guanine quartets. These structures have a low tolerance for non-guanine quartets which are rather formed by virtue of the docking platform provided by neighboring G-quartets. Few exceptions have been found. We demonstrate here that 8-amino guanine (R, Fig. 1) strongly accelerates quadruplex formation, which makes this nucleobase the most attractive replacement for guanine in the context of parallel quadruplexes.

We recently analysed the impact of a number of base analogs on parallel-stranded, tetramolecular quadruplex formation. 1 Most guanine substitutions are tolerated at best and are only formed thanks to the docking platform provided by neighboring G-quartets. Exceptions to this “guanine only” rule are guanine analogs modified at position 8. 1–3 Given the position of that substituent, which does not perturb the cyclic arrangement of H-bonds but favors the syn sugar base conformation, we wondered what would be the effect of other substituents at this position. 8-Amino guanine was previously found to stabilize triplexes 4 but destabilize quadruplexes 5 when inserted in the position 1 of an intramolecular quadruplex.

Following suggestions derived from preliminary theoretical calculations we decided to evaluate the effect of 8-amino guanine when inserted in the internal region of a parallel G-quadruplex.

Evidence for quadruplex formation with 8-amino G-containing sequences. Electrophoresis was used to evidence quadruplex formation (Fig. 2A). In native acrylamide gels, migration of the unmodified G4-DNA structure is characterized by a slight migration delay as compared to single-strands. By replacing each G in d(TG4T) or d(TG5T) sequences with R, we observed different migrations for structured and single-stranded forms.

Circular dichroism (CD) provided additional evidence for modified G4-DNA formation. CD spectra of all sequences are characterised by a negative peak at 242 nm, and a positive peak at 263 nm (Fig. 2B), referenced as “type 1 spectrum” in the literature. Isothermal difference spectra (IDS) of both modified and unmodified sequences were also characteristic for G4-DNA formation with two positive peaks at 240 and 273 nm, and two negative peaks at 260 and 295 nm. The shapes of these IDS confirm that all sequences form quadruplexes.

NMR spectra of all unmodified and modified sequences were consistent with quadruplex formation (see ESI). In all cases, guanine imino protons were observed around 11.0–11.5 ppm. These signals were still observed at very high temperatures (> 75 °C). In general, the modified oligonucleotides exhibited broader signals than the unmodified ones at low temperature (5 °C). These signals became narrower at higher temperatures. The spectra of [d(TGGRGGT)]₄ was similar to unmodified [d(TGGGGGT)]₄ and the non-exchangeable resonances could be assigned. On the other hand, there were significant changes in the NMR spectra of [d(TRGGGGT)]₄ and [d(TGGGGGT)]₄. Similar effects were observed in the spectra of [d(TRGGGT)]₄ and [d(TGGGGT)]₄. The exceeding

8-Amino guanine accelerates tetramolecular G-quadruplex formation†‡

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Fig. 1 8-Amino guanine (R, left) and schematic representation of the putative “R-tetrad” formed by the auto association of 4 R (right).
number of signals, probably due to the presence of minor species, make difficult the assignment of the NMR spectra of these modified quadruplexes. As in previously studied structures containing 8-amino guanines, the signals of the amino protons at position 8 were not observed in the NMR spectra. As a consequence, the conformation of their glycosidic angles could not be determined but, in all sequences, the medium or weak intensities of the H8-H1 NOEs indicated that the glycosidic angles of the unmodified guanines were anti. NMR data were consistent with quantum mechanical calculations which suggested no difference in the syn/anti free energy difference between guanine and 8-amino guanine (ΔG_{syn/anti} (R-G) = 0.1 kcal mol^{-1}; see supplementary Methods). There also was a clear agreement between NMR and MD simulations, which strongly suggest that minimum changes were due to the presence of 8-amino guanine in the quadruplex.

Association of G4 DNA. Association of a G4-DNA, which may be followed by absorbance, is generally slow and depends on temperature, oligonucleotide concentration, and nature and concentration of the monovalent cation.\cite{1,6–9} Replacement of a single G by 8-amino guanine (R) at various positions of d(TG₃T) or d(TG₄T) has been tested in this study and results are presented in Table 1.

In summary:

(1) As for both unmodified sequences, association rate of any modified sequence depends on the nature of the cation type: \( k_{\text{on}}(K^+) > k_{\text{on}}(Na^+) > k_{\text{on}}(NH_4^+) \) and its concentration \( k_{\text{on}}(K^+ \text{ 0.11M}) > k_{\text{on}}(K^+ \text{ 0.05M}) > k_{\text{on}}(K^+ \text{ 0.01M}) \).

(2) d(TG₃T) always leads to faster associations than d(TG₄T), and similar results are found for modified sequences. For example, \( k_{\text{on}}(\text{d}[\text{TRGGGTT}]_4) > k_{\text{on}}(\text{d}[\text{TRGGGT}]_4) \).

(3) Replacement at the 5’ end is more favorable than at the 3’ end, while insertion of R at central position hinders the association: \( k_{\text{on}}(\text{d}[\text{TRGGGTT}]_4) > k_{\text{on}}(\text{d}[\text{TGGGGTT}]_4) \). A similar 5’/3’ asymmetry was found for the shorter sequences.

(4) When considering the 5’ position, R leads to the fastest association kinetics among all tested bases, including guanine and 8-bromo guanine (X). Fig. 3 presents the kinetics of formation of the control sequence compared to the modified oligonucleotides. It illustrates the rate increased observed with the d(TRGGGTT) sequence.

![Fig. 2](image_url)  
Fig. 2 (A) Tetramolecular quadruplex formation is correlated to the apparition of slow-migrating bands (full triangle) as compared to migration of single-strands (open triangle). Results obtained in sodium conditions with TGG₃T sequences bearing an R at positions 1, 2, 3 or 4. Samples “+” were preincubated during 72 h at 4 °C at 100 μM strand concentration in a 10 mM lithium cacodylate pH 7.2 buffer containing 110 mM NaCl. In parallel, samples “-” were incubated under the same conditions omitting cations. (B) CD spectra in 10 mM lithium cacodylate pH 7.2 buffer containing 110 mM NH₄Cl.

![Fig. 3](image_url)  
Fig. 3 Accelerated quadruplex formation with 8-amino guanine. Acceleration of the association by replacing the first G at 5’ end on TG₃T by 8-bromo guanine (X) or 8-amino guanine (R). The three oligonucleotides have been used at the same concentration: (8.7 ± 0.15) μM. Association was followed by absorbance at (3.9 ± 0.3) °C in a 10 mM lithium cacodylate pH 7.2 buffer with 10 mM KCl + 100 mM LiCl.

### Table 1 \( k_{\text{on}} \) and \( T_{1/2} \) for TG₃T and TG₄T variants\(^{a,b}\)

<table>
<thead>
<tr>
<th>Sequences</th>
<th>( k_{\text{on}}/M^{-3}s^{-1})</th>
<th>( T_{1/2}(\text{C}°) )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( K^+ )</td>
<td>( Na^+ )</td>
</tr>
<tr>
<td>TGGGGGGT</td>
<td>2.3 × 10^{12}</td>
<td>9.8 × 10^{9}</td>
</tr>
<tr>
<td>TXGGGGT</td>
<td>1.2 × 10^{13}</td>
<td>1.1 × 10^{11}</td>
</tr>
<tr>
<td>TRGGGGT</td>
<td>Fast</td>
<td>Fast</td>
</tr>
<tr>
<td>TGGGGKT</td>
<td>6.9 × 10^{10}</td>
<td>8.7 × 10^{8}</td>
</tr>
<tr>
<td>TGGGGKT</td>
<td>3.9 × 10^{11}</td>
<td>6.5 × 10^{9}</td>
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<td>1.8 × 10^{10}</td>
<td>3.5 × 10^{8}</td>
</tr>
<tr>
<td>TRGGGGT</td>
<td>4.4 × 10^{11}</td>
<td>1.7 × 10^{10}</td>
</tr>
<tr>
<td>TGGGGT</td>
<td>2.1 × 10^{10}</td>
<td>3.6 × 10^{9}</td>
</tr>
</tbody>
</table>

\(^{a}\) X and R correspond to 8-bromo-guanine and 8-amino guanine, respectively. 
\(^{b}\) The association rate (at 4 °C) is provided in M^{-3}s^{-1} (assuming fourth order association kinetics\(^{1,6};\) see a recent discussion in ref. 10) in a 10 mM lithium cacodylate pH 7.2 buffer supplemented with 110 mM KCl, NaCl or LiCl. Nd: not done. Fast: formation of the quadruplex is too fast at 110 mM KCl to be followed under these ionic conditions even at the lowest practical strand concentration. At 10 mM KCl, association of TRGGGGT is 7-fold and 63-fold faster than TXGGGGT and TGGGGGT, respectively (not shown). \(^{\dagger}\) T_{1/2} in °C correspond to non-reversible apparent melting temperatures and depend on the temperature gradient (0.5 °C min^{-1} here); >90: no significant denaturation occurred during the melting experiment.
Methods

The lack of destabilization (or even the gain of stability) of quadruplexes by 8-amino guanine insertion in the middle of the structure seems in disagreement with previous theoretical and experimental studies, which found destabilization of the structure when \( R \) replaced one \( G \) in a two-step intramolecular quadruplex.\(^5\) In order to solve this apparent paradox, we performed MD/TI calculations similar to those in ref. 5, but now considering none, one, or four \( G \) to \( R \) substitutions at the 2nd step of the \([dGGGG])_4\) quadruplex (see supplementary Methods).\(^\dagger\) Results showed an important gain of stability (around 2.4 kcal mol\(^{-1}\)) when one guanine is substituted by one 8-amino guanine, the difference being enlarged to 9.5 kcal mol\(^{-1}\) when the entire 2nd step is changed to 8-amino guanines, in agreement with experimental measures and in opposition to what was found previously in the analysis of external G to R changes. Theoretical calculations therefore support the marked sequence-dependent effect of the G to R substitution and the fact that the introduction of the modified nucleotide in the interior of the G-quadruplex can stabilize the structure.

Fig. 4 Effect of \( R \) on G4 formation and thermal stability. Melting profiles of G4-DNA prepared in NaCl 110 mM, based on \( d(TG\_4T) \) sequences and modified at each positions with \( R \).

Notes and references