The formation pathway of tetramolecular G-quadruplexes

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ABSTRACT

Oligonucleotides containing guanosine stretches associate into tetrameric structures stabilized by monovalent ions. In order to describe the sequence of reactions leading to association of four identical strands, we measured by NMR the formation and dissociation rates of (TGNnT)₄ quadruplexes (n = 3–6), their dissociation constants and the reaction orders for quadruplex formation. The quadruplex formation rates increase with the salt concentration but weakly depend on the nature (K⁺, Na⁺ or Li⁺) of the counter ions. The activation energies for quadruplex formation are negative. The quadruplex lifetimes strongly increase with the G-tract length and are much more longer in K⁺ solution than in Na⁺ or Li⁺ solutions. The reaction order for quadruplex formation is 3 in 0.125 M KCl and 4 in LiCl solutions. The kinetics measurements suggest that quadruplex formation proceeds step by step via sequential strand association into duplex and triplex intermediate species. Triplex formation is rate limiting in 0.125 M KCl solution. In LiCl, each step of the association process depends on the strand concentration. Parallel reactions to formation of the fully matched canonical quadruplex may result in kinetically trapped mismatched quadruplexes making the canonical quadruplex practically inaccessible in particular at low temperature in KCl solution.

INTRODUCTION

Two nucleic acid families can form tetrameric structures. The oligonucleotides containing stretches of at least two adjacent cytidines associate into a structure, the i-motif, formed by two parallel duplexes held by hemiprotonated C–C⁺+ pairs and intercalated in each other (1). The G rich oligomers form four stranded structures maintained by stacked G-quartets (2–4). The G-quadruplexes and the i-motif have been the object of detailed structural studies (5), but the sequence of the reactions leading to the association of four strands into a tetrameric structure is not clear. The present article reports a kinetics study of the formation and dissociation of (TGNnT)₄ quadruplexes (n = 3–6) in KCl, NaCl and LiCl solutions and of (rUG4U)₄ in NaCl solution, performed with the goal of describing their formation pathway. An equivalent study of the i-motif formation pathway is in preparation.

The (TGNnT)₄ quadruplexes are formed by four parallel strands bound together by n layers of G quartets (4). They include n – 1 monovalent cations bound between the G-quartets in the central cavity. This structure, that is readily identifiable by its NMR spectrum (6–8), is considered as the thermodynamically stable strand assembly. It will be designated as the ‘canonical quadruplex’.

It is unrealistic to imagine that quadruplex formation could result from the simultaneous collision of four strands. Quadruplex formation proceeds without doubt by step-wise strand association into elusive duplex or/and triplex structures. Nevertheless, there is no experimental evidence about the presence of such intermediate species. Kinetics experiments, including the measure of the quadruplex formation and dissociation rates, of the reaction order and equilibrium measurements of the quadruplex dissociation constant may provide information leading to the description of the association pathway.

Using gel filtration chromatography (9) and optical methods (10–15), previous kinetics studies have shown that strand association and quadruplex dissociation is extremely slow. A model for G-quadruplex formation involving association of two preformed duplexes was proposed on the basis of gel filtration chromatography experiments (9). NMR provides, by contrast with these methods, direct information about the nature of the multimers formed. Nevertheless, NMR has never been used to investigate the G-quadruplex formation kinetics. It will be shown that canonical quadruplex formation may go together with faster parallel reactions resulting in the apparition of quadruplex structures that are kinetically trapped in the early phase of the association reaction. The formation of the canonical quadruplex may be hindered

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in that case during periods exceeding any possibility of kinetics investigation. A rigorous control of the nature of the multimers formed is therefore a prerequisite condition to G-quadruplex kinetics investigation. For each oligonucleotide examined, we selected the experimental conditions (salt concentration, temperature) in which strand association leads to the formation of a single species: the canonical G-quadruplex. The description of the G-quadruplex formation pathway derived from these experiments is distinct of that initially proposed (9). Nevertheless, the data provided by the previous kinetics studies were a great help in the conception of the present study by allowing us to focus the NMR investigations in directions already explored.

MATERIALS AND METHODS

Oligonucleotides and NMR samples

The oligonucleotides were synthesized on a 10 μM scale and purified by chromatography on a DEAE column as previously reported (16). Depending on the cation used in the kinetics experiments, the deoxy oligonucleotides were dissolved after purification in 0.1 M Li⁺, Na⁺ or K⁺ hydroxide solutions in order to destroy any structure formed. The RNA sample was dissociated by heating at 95°C during 5 min. Afterwards, the oligonucleotides were dialyzed, first against water, then against 0.1 M LiCl, NaCl or KCl solutions and repeatedly against water. At last, the oligonucleotides were lyophilized and dissolved in water. The solution pH was adjusted to 6.5 with HCl and Li⁺, Na⁺ or K⁺ hydroxide. The strand concentration was determined using the A 260 values provided by Mergny et al. (12). The NMR samples were usually buffered by Li⁺, Na⁺ or K⁺ phosphate 20 mM, pH 6.5. For the experiments performed at low salt concentration, the phosphate concentration was reduced to 2 mM. The monovalent cation concentration was adjusted by addition of 1 M Li⁺, Na⁺ or KCl solutions. Two microliters of a solution containing 100 mM dimethyl sulfoxide (DMSO) and Li⁺, Na⁺ or K⁺ acetate (50 mM) was added to each NMR sample. The chemical shift of the methyl protons of acetate provided a marker for pH determination (17). The DMSO NMR peak was used as an internal marker allowing normalization of spectra collected during the kinetics experiments. Ethylene diamine tetra acetic acid, Na⁺ form, was added to the RNA samples at the final concentration of 0.5 mM.

Association and dissociation rate measurements

To measure the quadruplex formation rates, the oligonucleotide solutions were melted at 100°C in the NMR tube and rapidly inserted into the NMR probe (17). The quadruplex dissociation time at 100°C can be extremely long in the presence of salt. In that case, the monovalent salt was added into the NMR tube immediately after melting. To measure the quadruplex dissociation rates, a small volume of a concentrated fully tetrameric oligonucleotide solution was diluted in the NMR tube immediately before the beginning of the experiment. The dilution buffer was degassed in order to avoid bubble formation during the experiments performed at high temperature. The NMR probe cannot be used at temperature higher than 80°C. In that case, the NMR samples were stored in a thermostated incubator and the monomer and quadruplex fractions were measured on spectra recorded at 75°C at different times during incubation.

The NMR experiments were performed on a 500 MHz Varian Inova spectrometer using the jump and return sequence for water suppression (18). The dead time to start the kinetics measurements was shorter than 3 min. Typically, the kinetic experiments were run overnight or during a weekend. Afterward, the NMR tube was sealed with parafilm and stored at controlled temperature in a thermostated incubator. Additional kinetics measurements were performed on the same sample after several days or weeks.

The tetramer and monomer fractions were determined from the area of several markers. The monomer and tetramer T methyl peaks provide the best markers due to their larger intensity. At high temperature, the well-resolved G NIH peaks were used as a marker of the quadruplex fractions. Depending on temperature and salt concentration, the T (H6) protons and several G (H8) protons belonging to the monomer and to the quadruplex were also used (Figure 1).

Determination of the reaction order of G tetramer formation

The reaction order for quadruplex (Q) formation is the value of the exponent of the monomer concentration (M) in the rate equation: d(Q(t))/dt = k[M(t)]^order.

The half reaction time, tasso, M0 for strand association into a multimeric structure depends on a rate constant, k, on the initial strand concentration [M0] and on the reaction order:

\[ t_{asso, M0} = \frac{1}{k[M0]^{order-1}} \]

Figure 1. Aromatic and methyl proton region of spectra recorded during the association kinetics of TG3T. The spectra were collected at different times right after melting of the oligonucleotide. The NMR peaks of the monomer are labeled ‘c’ and those of the quadruplex are labeled by the residue number according to Jin et al. (8). The line labeled by a star is that of acetate, a marker used to control the solution pH. The aromatic and methyl proton regions of the NMR spectrum show several well-resolved lines providing markers to measure the fraction of each species as a function of the time. The small chemical shifts observed for several NMR peaks of the monomer are indicated by dotted lines. They reflect a fast exchange situation between the monomer and an intermediate species, presumably (TG3T)2 (see text). Solution conditions: (TG3T) = 1.5 mM, T = 0°C, [K⁺] = 10 mM.
The quadruplex dissociation constant was determined from the slope, \(-(\text{order} - 1)\), of the log–log plot of the half reaction time versus the oligonucleotide concentration. The error on \(\tau_{\text{asso,M0}}\) values evaluated by using different markers of the NMR spectra is \(\sim 15\%\). We estimate that the error on the reaction order is comparable.

### The quadruplex dissociation constant

The quadruplex dissociation constant in an oligonucleotide solution at concentration \([M0]\) may be calculated from the monomer fraction at equilibrium, \(\alpha_{\text{eq}}\), by:

\[
K_{\text{dis}} = \frac{4\alpha_{\text{eq}}^2[M0]^3}{(1 - \alpha_{\text{eq}})}.
\]

The dimension of the dissociation constant is \(M^3\). It is more intelligible to express the quadruplex stability by \(Fi\), the monomer concentration in equilibrium with the quadruplex when \(\alpha_{\text{eq}} = 0.5\). It follows from this definition that the oligonucleotide concentration of a G-quadruplex solution at the melting temperature is \(2 \cdot Fi\). \(Fi\), that will be designated as the reduced dissociation constant is related to the quadruplex dissociation constant by:

\[
Fi = \left(\frac{K_{\text{dis}}}{4}\right)^{(1/3)}.
\]

The coherence of the tetramer and monomer equilibrium concentrations measured at different oligonucleotide concentrations was systematically controlled by checking that the tetramer concentration increases as the power of four of the monomer concentration.

### Gel filtration chromatography

The stoichiometry of the structures formed by \(TG^nT\) association was controlled at room temperature by high-pressure chromatography using a Synchropack GPC 100 column in 0.1 M Na or KCl elution solutions. The column was calibrated using \(TG^nT\) monomers, deoxyduplexes containing from 6 to 16 bp and a tRNA sample.

# RESULTS

### Association kinetics

For each set of experimental condition (temperature, oligonucleotide concentration, nature and concentration of the monovalent cation) the monomer and quadruplex concentrations were measured as a function of the time and the tetramer reduced dissociation constant was determined from the evaluation of the concentration of each species at equilibrium.

The results related to \(TG^nT\) association are sorted in two sub-sections according to a simple criterion: the presence or the absence during the association kinetics of any multimeric species different from the canonical quadruplex. In the first section, we present the experiments showing the formation of a single species: the canonical quadruplex. In that case, the half reaction times measured for monomer and quadruplex evolutions are identical. In the second sub section, we describe several experiments showing, or suggesting the formation of kinetically trapped quadruplexes distinct of the canonical quadruplexes. In that case, the evolution of the monomer concentration is much faster than that of the canonical quadruplex.

### Formation kinetics of canonical \((TG^nT)_4\) quadruplexes

**TG3T.** Representative spectra collected during the formation of \((TG3T)_4\) in a 1.5 mM \(TG3T\) solution initially monomeric are displayed in Figure 1. The spectra show only two species: the monomer, identified by comparison with the spectrum of a dilute \(TG3T\) sample at 70°C and the canonical tetramer. The sum of the monomer and tetramer fractions measured during the kinetics is close to one (Figure 2). This, together with the identity of the half reaction time of each species, 40 ± 5 min, establishes that formation of any other multimer is negligible. At equilibrium, the free strand proportion, \(\alpha_{\text{eq}} = 0.1\), corresponds the reduced dissociation constant \(Fi = 7.2 \times 10^{-3}\) M. Figure 2 shows that the half association times, \(\tau_{\text{asso}}\), measured versus the oligonucleotide concentration in 0.125 and 10 mM KCl solutions at 0°C vary as the power of \(-2\) of the oligonucleotide concentration and thus establishes, Equation (1), that the reaction order for \((TG3T)_4\) formation is 3. The half formation time of \((TG3T)_4\) increases between 0 and 37°C with an activation energy of \(-36.2\) kJ/mol (Figure S1). The reaction order is 3 in all this temperature range (Figure 3). The half formation time of \((TG3T)_4\) decreases with the KCl concentration approximately as \([\text{KCl}]^{-1.8}\) (Figure S2). The reaction order for \((TG3T)_4\) formation in 37 mM KCl at 21°C is 3.2 ± 0.2.

In 0.125 M NaCl, the half formation time of \((TG3T)_4\) at 0°C is about three times shorter than that measured in 0.125 M KCl. Due to the poor stability of \((TG3T)_4\) in NaCl solution \((Fi = 3 \times 10^{-4}\) at 0°C) and to the fast association rate at high oligonucleotide concentration, the reaction order in NaCl solution, 3.5 ± 1, could not be determined with accuracy.

**TG4T.** In 0.125 M KCl solution, the formation rate of \((TG4T)_4\) measured versus the oligonucleotide concentration between 0 and 72°C establishes that the reaction order is 3 (Figure S3). By contrast, in 5 mM KCl at 37°C, \(\tau_{\text{asso}}\) vary as the power of \(-3\) of the oligonucleotide concentration indicating that the reaction order is 4 (Figure S3).

In 0.125 M LiCl, the half formation time of \((TG4T)_4\) is longer that in KCl solution (Figure 4). The variation of \(\tau_{\text{asso}}\) versus the oligonucleotide concentration measured between 0 and 37°C shows that the reaction order for \((TG4T)_4\) formation is 4 (Figures 3 and S3).

In 0.125 M NaCl, the apparent reaction order increases with temperature from 3.2 ± 0.1 at 0°C up to 3.6 ± 0.1 at 50°C (Figure 3). It depends also on the NaCl concentration. At 21°C, the reaction order increases from 3.2 in 600 mM NaCl up to 4 in 37.5 mM NaCl (Figure 3). The half formation time of \((TG4T)_4\) decreases with the NaCl concentration similarly to that of \((TG3T)_4\) versus the KCl concentration (Figure S2).

**TG5T.** The spectra collected during the association kinetics of \(TG5T\) at 42 and 60°C in 0.125 M NaCl or LiCl shows also that the monomer and canonical
quadruplex fractions vary simultaneously. The reaction order for (TG5T)₄ formation at 42°C is 3 in NaCl and 4 in LiCl solution (Figure 3). The half formation times of (TG5T)₄ and (TG4T)₄ measured in 10 mM oligonucleotide solutions at 60°C in 0.125 M NaCl are similar (Figures 4 and S1) but the activation energy for (TG5T)₄ formation is about twice that measured for (TG4T)₄ (Table 1).

Parallel formation of mismatched and canonical quadruplexes. In a number of cases, we observed that one or several multimers are formed more rapidly than the canonical duplex. The formation of such structures is favored at low temperature, in the presence of KCl and by the G-tract length of the oligonucleotide sequences (Scheme 1).

A clear example is shown on the spectra recorded during the association kinetics of TG5T at 42°C in 0.125 M NaCl. In the early stage of the experiment, the spectra show the accumulation of a species, Qₓ, distinct of the canonical tetramer, Q (Figure 5). Size exclusion chromatography experiments performed on samples taken off the NMR tube at different times during the association kinetics show that the Q and Qₓ species are eluted as a single peak with the elution time expected for a tetramer. The strand arrangement in the Qₓ tetramer could not be determined due to its limited lifetime and poor spectral resolution. Qₓ may be a quadruplex with one or several slipped strands or with anti-parallel strands.

In 10⁻⁴ M TG5T solution, the half association time of the monomeric species is ~500 min. The Qₓ fraction at equilibrium, fₓ = 0.25, yields the reduced dissociation constant: Fᵣ = 7.2 x 10⁻⁵ M. Right panel: half reaction time for (TG3T)₄ formation versus the oligonucleotide concentration at 0°C in 0.125 M (black squares) and 10 mM (black circles) KCl. The lines drawn through the data points show that the half reaction time increases as power of ~3 of the oligonucleotide concentration and therefore establishes that the reaction order is 3.

rUG4U. The reaction order for (rUG4U)₄ formation in 0.125 M NaCl is 3.4 at 42°C and 3.8 at 60°C (Figure 3).

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order for Q and Qx formation is 3 as in the case of (TG3T)₄ and (TG4T)₄ at the same KCl concentration. A model involving two parallel reactions accounts for the time evolution of each species in 10⁻⁴ to 10⁻³ TG5T solutions.

\[
<---k_{on1} \kappa_{on2}--> \quad Qx \quad \text{monomer} \quad Q
\]

\[k_{off1} \quad <---k_{off2}\]

The fits displayed in Figure 5 were computed with the formation rates \(k_{on1} = 1/160 \text{ min}^{-1}\) for Qx and \(k_{on2} = 1/700 \text{ min}^{-1}\) for Q and with the Qx dissociation rate \(k_{off1} = 1/870 \text{ min}^{-1}\). The predominance of the canonical quadruplex at equilibrium is due to its extremely long lifetime: \(1/k_{off2} \geq 1 \text{ year}\). According to Equation (2), the Qx fraction decreases versus time with a rate constant \(k = k_{off1} \kappa_{on2}/(k_{on1} + k_{on2})\). It may be noted that subsequently to Qx dissociation, the back reaction towards Qx is faster than Q formation, i.e. \(k_{on1} > k_{on2}\). For this reason the Qx fraction decreases with a time constant, \(~4600 \text{ min}^{-1}\), longer than the Qx lifetime (Figure 5). This is a typical example of kinetics trapping, a phenomenon observed in parallels equilibrium when the formation rate of the less stable reaction product (Qx in this example) is faster than that of the thermodynamically stable species (Q in this example).

The NMR spectra recorded during the association kinetics of TG4T in 0.125 M KCl at temperature lower than 20°C, of rUG4U in 0.125 M NaCl at temperature lower than 42°C and of TG6T in 0.125 M Li⁺, Na⁺ and KCl solutions suggest also the formation of species distinct of the canonical quadruplex (Scheme 1). In these experiments, the good resolution of the monomer markers allowed an accurate determination of the monomer half association times, but it was impossible to determine the formation rates of the multimers due to the poor spectral resolution.

In 0.125 M KCl, the half association time of TG4T in 10⁻⁴ M solution is 5 min at 0°C (Figure 4). The spectra recorded during several weeks, show extremely broad multimer peaks and suggest that the equilibrium is reached. Gel filtration chromatography of aliquots removed from the NMR tube indicates that the structure(s) formed is (are) tetrameric. Nevertheless, after incubation at 42°C during 1 day and slow cooling.

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**Figure 4.** Effect of temperature on the formation and dissociation half lifetime and on the reduced dissociation constant of (TG4T)₄ in 0.125 M KCl, NaCl and LiCl. Upper panels: quadruplex lifetimes (triangles). Half formation times of the canonical quadruplex (open squares) and half association times of the TG4T monomer (black circles) measured in 10⁻⁴ M oligonucleotide solutions or extrapolated at 10⁻³ M from measures at higher oligonucleotide concentrations. The half lifetimes for canonical quadruplex formation and for monomer association are identical in the range of temperature where the formation of mismatched quadruplex is negligible (Table 1). Lower panel: reduced equilibrium constants derived from the oligonucleotide concentrations. The half lifetimes for canonical quadruplex formation and for monomer association are identical in the range of temperature where the formation of mismatched quadruplex is negligible (Table 1). Lower panel: reduced equilibrium constants derived from the monomer concentration at equilibrium (open circle). The crosses indicate the reduced equilibrium constants computed from the association and dissociation rates used in the numerical simulations.

**Table 1.** Kinetics and equilibrium parameters for G-quadruplex formation and dissociation in 0.125 M salt

<table>
<thead>
<tr>
<th>Sequence</th>
<th>Salt</th>
<th>(A_{asso} (\text{min}))</th>
<th>(E_{asso} (\text{kJ/M}))</th>
<th>(A_{disso} (\text{min}))</th>
<th>(E_{disso} (\text{kJ/M}))</th>
<th>(Ft_{42} (\text{M}))</th>
<th>(Tm (\text{°C}))</th>
<th>(t_{disso}, T_e (\text{days}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>TG3T</td>
<td>K</td>
<td>(6.2 \times 10^{-10})</td>
<td>(-36.2)</td>
<td>(6.7 \times 10^{29})</td>
<td>190.7</td>
<td>(1.2 \times 10^{-3})</td>
<td>28 ± 3</td>
<td>2 ± 1</td>
</tr>
<tr>
<td>TG4T</td>
<td>K</td>
<td>(1.0^{-12})</td>
<td>(-93.1)</td>
<td>(2.8 \times 10^{28})</td>
<td>220.3</td>
<td>(1.4 \times 10^{-6})</td>
<td>69 ± 4</td>
<td>112 ± 49</td>
</tr>
<tr>
<td>TG4T</td>
<td>Na</td>
<td>(1.9 \times 10^{-12})</td>
<td>(-50.4)</td>
<td>(4.8 \times 10^{26})</td>
<td>177.5</td>
<td>(3.5 \times 10^{-4})</td>
<td>34 ± 4</td>
<td>2 ± 1</td>
</tr>
<tr>
<td>TG4T</td>
<td>Li</td>
<td>(8.3 \times 10^{-13})</td>
<td>(-44.9)</td>
<td>(1.4 \times 10^{28})</td>
<td>185.9</td>
<td>(5.4 \times 10^{-4})</td>
<td>24 ± 6</td>
<td>34 ± 22</td>
</tr>
<tr>
<td>TG5T</td>
<td>Na</td>
<td>(6.7 \times 10^{-20})</td>
<td>(-95.6)</td>
<td>(3.0 \times 10^{20})</td>
<td>365.6</td>
<td>(3.1 \times 10^{-7})</td>
<td>73 ± 4</td>
<td>38 ± 22</td>
</tr>
<tr>
<td>rUG4U</td>
<td>Na</td>
<td>(1.6 \times 10^{-30})</td>
<td>(-163.2)</td>
<td>(2.6 \times 10^{20})</td>
<td>297</td>
<td>(6 \times 10^{-7})</td>
<td>71 ± 5</td>
<td>50 ± 30</td>
</tr>
</tbody>
</table>

The quadruplex formation rates in 0.1 mM oligonucleotide solution are \(k_{asso,10^2} = A_{asso} \exp(-E_{asso}/RT)\) and the concentration independent quadruplex dissociation rate \(k_{disso} = A_{disso} \exp(-E_{disso}/RT)\). \(E_{asso}\) and \(E_{disso}\) are the activation energies for strand association and quadruplex dissociation. \(Ft_{42}\) is the reduced dissociation constant at 42°C. \(Tm\) is the quadruplex equilibrium melting temperature in 10⁻³ M oligonucleotide solution (see text). \(t_{disso}, T_e\) is the quadruplex dissociation time at the melting temperature.
(about 1°C/h) at 4°C, one observes a well-resolved spectrum that may be assigned to the canonical tetramer, by comparison with published NMR spectra (6,7). This indicates that TG4T association at 0°C forms mismatched tetrameric structures that are kinetically trapped during extremely long periods. Dissociation of these structures at 42°C allows the formation of the canonical tetramer during the slow cooling procedure. The slope change observed around 20°C on the plot of the half association time of TG4T versus temperature (Figure 4) is likely the consequence of the faster formation rate of mismatched structures.

The association kinetics of TG6T in 0.125 M NaCl at 42°C is about 100 times faster than that of TG4T, but the NMR spectra recorded during TG6T association show the formation of at least three species that are still present after incubation during 4 weeks. In that case again, gel filtration experiments show that the species formed are tetrameric. After heating at 100°C and slow cooling to room temperature (about 1°C/h), one observes a well-resolved spectrum indicating the formation of a single symmetrical quadruplex, presumably the canonical species. In 0.125 M KCl, the same annealing procedure increases the proportion of one of the quadruplexes formed.

The observations reported in this sub section show that association of TGN4T oligonucleotides may result in the formation of mismatched quadruplexes that are kinetically trapped. The length of the G tract favors the formation of these structures. They are preferentially formed at low temperature in the presence of KCl (Scheme 1).

Proton shifts during TGN4T association

The spectra collected as a function of the time during TG3T and TG4T association at 0°C show that several NMR peaks of the monomer exhibit systematic chemical shifts simultaneously with the formation of mismatched quadruplexes. The shift amplitude is modest. The largest, 0.015 p.p.m., is always that of the low field T CH3 peak (1.73 p.p.m. on the TG3T spectra displayed in Figure 1). The shift amplitude decreases when the temperature increases and becomes negligible above 20°C. The shifts measured in 0.125 M KCl and NaCl are comparable. Using experiments performed versus time to follow the formation kinetics of TG4T in 40 μM to 1 mM oligonucleotide solutions, we measured, on all the spectra recorded versus time, the monomer T CH3 frequency and the monomer concentration. The plot of the T CH3 frequency versus the monomer concentration shows that, independently of the oligonucleotide concentration, the spectra recorded with comparable free monomer concentrations are similarly shifted (Figure 6). This shows that the shifts are related to the monomer concentration and therefore this suggest that they result of a fast exchange equilibrium between the monomer and a short-lived intermediate species.
triangles) and 10 

solid lines were obtained by assuming dimer TCH3 frequencies of 1.5 monomer TCH3 frequency is estimated to 1.831 p.p.m. The computed monomer and an intermediate species, presumably (TG4T)2. The versus the monomer concentration suggests fast exchange between the 

The lifetime at 60 

addition of one residue to the G tract increases the 

of the counter ion and on the G-tract length (Figure 7). 

exponential function with a time constant independent of 

during dissociation was satisfactorily fitted by a mono 

Tetramer dissociation kinetics 

Tetramer dissociation opposes a one-order dissociation reaction to a multi-order formation reaction. Hence, the measured dissociation time depends on the oligonucleotide concentration and on the association and dissociation rate constants when dissociation at equilibrium is not achieved. In order to determine the quadruplex lifetimes, the dissociation kinetics experiments were always performed using solutions diluted in such a way that the quadruplex equilibrium fraction was <0.1. On this condition, the evolution of the tetramer concentrations during dissociation was satisfactorily fitted by a mono exponential function with a time constant independent of the oligonucleotide concentration. 

The quadruplex lifetimes depend strongly on the nature of the counter ion and on the G-tract length (Figure 7). At 60°C, the lifetime of (TG4T)4 in 0.125 M NaCl, 12 min, is comparable to that measured in LiCl solution, 8 min. It is much more longer in 0.125 M KCl, about 2 years, as shown by the extrapolation of the lifetimes measured between 72 and 100°C. The activations energies measured for tetramer dissociation in 0.125 M K+ and Na+ and LiCl solutions are listed in Table 1. The comparison of the lifetimes of (TG4T)4 and (TG3T)4 in 0.125 M KCl or of (TG4T)4 and (TG5T)4 in 0.125 M NaCl shows that addition of one residue to the G tract increases the tetramer lifetime by 5 to 6 magnitude orders (Figure 7). The lifetime at 60°C of (rUG4U)4 is about 10^2 times longer than that of the homologous DNA tetramer and its activation energy, 297 ± 30 kJ/M, is much higher (Table 1). 

We also examined the effect of the NaCl concentration on the dissociation kinetics of (TG4T)4 and the effect of KCl on (TG3T)4 and (TG4T)4 dissociation. In agreement with previous observations (12), we observe that the lifetime of (TG4T)4 measured at 58°C in NaCl solutions ranging from 5 to 500 mM is independent of the NaCl concentration. By contrast, in KCl solution the lifetimes of (TG3T)4 (measured at 28°C) and of (TG4T)4 (measured at 86°C) decrease as the KCl concentration. Nevertheless, the quadruplex lifetimes measured at the lowest KCl concentration examined, 2 mM, are several magnitude orders longer than those measured in NaCl solution.

Tetramer stability 

The reduced dissociation constants, Fi, displayed in Figures 4 and S1 were obtained from the free and bound strand concentrations measured on the last spectra recorded during the kinetics measurements. Table 1 shows that addition of one residue to the G tract decreases Fi by about 3 magnitude orders in NaCl as in KCl solution.

Fi measurements yield redundant information that may be independently computed from the dissociation and association rates. The comparison of the Fi values measured at equilibrium with those derived from the kinetics parameters provides a rigorous control of the coherence of the kinetics and equilibrium measurements. The G-quadruplex melting temperature, T_{m,M0}, in an oligonucleotide solution of concentration [M0] is the temperature for which the half association time, \tau_{asso,M0} is equal to the concentration independent dissociation half lifetime, \tau_{diss}. Furthermore, according to the Fi definition, the G-quadruplex reduced dissociation constant at the melting temperature is Fi = [M0]/2.

The quadruplex melting temperatures in 10^-4 M oligonucleotide solution, T_{m,10^-4}, are determined in the upper panels of Figures 4 and S1 by the interception of the plots of \tau_{asso,10^-4} and \tau_{diss} versus temperature. The lower
panels of Figures 4 and S1 show that the $F_i$ values at temperature $T_{m,10^{-4}}$ are close to $10^{-4}/2$ and thus establishes the consistency of the equilibrium and kinetic measurements.

**DISCUSSION**

The slow association and dissociation kinetics of the G-quadruplexes has been noticed since the very beginning of their discovery and repeatedly discussed and analyzed since that time (9–15). In fact, one should not be surprised by the slow formation rates of these structures. It is hardly conceivable that association of four dilute strands via a process involving unstable intermediate species into a structure requiring correct strand orientation and alignment could be fast. More surprising are the extraordinarily long tetramer lifetimes that extrapolate to million of years at 0°C as in the case of TG4T in KCl solution (Figure 7) and the huge activation energies for tetramer dissociation (Table 1) indicating that a quasi-insurmountable energetic barrier must be overcome to dissociate the quadruplex.

The association kinetics of TG5T in KCl shows how the fast formation rate of a mismatched species and its relatively long lifetime can make practically inaccessible the thermodynamically stable canonical quadruplex. With the exception of TG3T, this process seems to be the general rule between 0 and 20°C in KCl solution. The characterization of the structure(s) formed by strand association is for this reason essential to the interpretation of the G-tetramer formation kinetics.

The association process of TGnT oligonucleotides into canonical quadruplex

We determined for each oligonucleotide of the TGnT series the experimental conditions (salt concentration, temperature) leading to the formation of the canonical tetramer (Scheme 1). The model discussed in this section relies on the parameters (formation and dissociation time constants, reduced dissociation constants and reaction orders) obtained from these experiments.

The reaction order is the key parameter to determine a reaction pathway. In summary, we find that the order for quadruplex formation depends on the nature of the counter ion. In 0.125 KCl, the formation order is always 3 and it is always 4 in LiCl solution. It depends on the monovalent ion concentration and temperature. In NaCl solution, the order is 3, or close to 3, in conditions favorable to multimer stability (i.e. low temperature and high salt concentration). It is close to four at low NaCl (Figure 3) or KCl concentrations (Figures 3 and S3).

The reaction order depends also on the G-tract length, as shown by the comparison of the reaction orders measured for (TG4T)$_4$ (order 3.5) and (TG5T)$_4$ (order 3) (Figure 3).

Two models accounting for quadruplex formation can be considered (19). The model initially proposed by Wyatt et al. (9) involves two steps: fast pre-equilibrium between monomer (M) and dimer (D) and duplex association into a quadruplex structure (Q).

\[
\begin{align*}
&k_{on1} \quad \text{step 1} \\
&2M \quad \text{D} \\
&k_{on2} \quad \text{step 2} \\
&2D \quad Q
\end{align*}
\]

Where, $k_{on1}$ and $k_{of}$ are the association and dissociation rates at each step. According to this model, the reaction orders for Q formation are four if both steps are limiting. If duplex association into quadruplex is much faster that duplex dissociation, step 2 should be non limiting, resulting in a reaction order of two. But this scheme is by no way compatible with the association order of 3 measured for tetramer formation in KCl solution.

The second model involves step-by-step strand association into duplex, triplex (T) and quadruplex structures.

\[
\begin{align*}
&k_{on1} \quad \text{step 1} \\
&2M \quad D \\
&k_{on2} \quad \text{step 2} \\
&M + D \quad T \\
&k_{on3} \quad \text{step 3} \\
&M + T \quad Q
\end{align*}
\]

According to this scheme, the reaction order for Q formation is four if the reactants are in rapid pre-equilibrium at every step but it may be smaller if one step is non-limiting. Step 1 is obviously in rapid pre-equilibrium, otherwise accumulation of the D species should be observed and the reaction order should be ≤2. If step 2 is non-limiting, accumulation of the triplex species during the early stage of the kinetics should result in the apparition of new NMR peaks. From the absence of evidence in favor of this assumption we conclude that only step 3 may be non-limiting. The model proposed for G-quartet formation rely on the assumption that step 3 is non limiting in the experimental condition for which the reaction order is 3, i.e. (TG4T)$_4$ formation in 0.125 M KCl, and limiting when the reaction order is four, i.e. (TG4T)$_4$ formation in LiCl solution.

The quadruplex formation rate across step 3 may be expressed by:

\[
k_3 = \frac{k_{on3}[M][T]}{(k_{of2} + k_{on3}[M][T])^p}, \quad \text{or} \quad k_3 = \frac{p[M]}{(1 + p[M])}, \quad \text{with} \quad p = k_{on3}/k_{of2}
\]

This expression shows immediately that step 3 is not limiting when M association to T is faster than T
and 4 in 5 mM KCl (Figure S3) may be similarly explained. At last, the difference observed in 0.125 M NaCl (Figure 3) between the formation orders of (TG4T)$_d$ (order 3.5) and (TG5)$_d$ (order 3) may reflect a longer lifetime of the intermediate triplex structure of TG5T.

The methyl shifts observed versus the free monomer concentration at 0°C (Figure 6) might be the consequence of the fast monomer-duplex pre-equilibrium at step 1. According to this assumption the methyl shift, δ, should depend on the monomer and dimer frequencies, δ$_1$ and δ$_2$ respectively, and on $f_{d1}$ the dimer fraction: 

$$\delta = (1-f_{d1})\delta_1 + (f_{d1})\delta_2.$$ 

Here, $f_{d1}$ and the dimer dissociation constant, $K_d = (1-f_{d1})/([monomer] f_{d1})^2$, may be estimated from the plot of δ versus [monomer] if one knows δ$_1$ and δ$_2$. δ$_1$ may be estimated to 1.831 p.p.m. from methyl frequency at the lowest monomer concentrations. The determination of δ$_2$ requires shift measurements at high monomer concentrations that are impracticable due to fast tetramer formation (Figure 2, right panel). To estimate the range of the $K_d$ values accounting for the measured shifts we tried different pairs of values for the δ$_2$ and $K_d$ parameters. The fits displayed on Figure 6 were obtained with δ$_2 = 1.6$ and 1.5 p.p.m. and dimer dissociation constants $K_d = 2 \times 10^{-3}$ and 6 x $10^{-3}$ M$^{-1}$.

Kinetic simulations performed, as described in Supplementary Data, by numerical integration are displayed in Figures 8 and S4.

Considering that the monomer and duplex are in fast pre-equilibrium, we can write:

$$k'_{on1} > 3M \quad T \quad (step 1 + step 2)$$

$$< --- --- >$$

$$M + T \quad Q \quad step 3$$

$$< --- --- >$$

$$k'_{off3}$$

The kinetics experiments provide three independent parameters: $\tau_{asso}$, $\tau_{disso}$ and the reaction order. It is therefore impossible to determine the four rates of (step 1 + step 2) and step 3. The rate constant $k'_{off3}$ was identified with the inverse of the quadruplex lifetime (Figure 4 and Table 1) and $k'_{on1}$ was varied to obtain a good agreement with the monomer and quadruplex fractions measured as a function of the time. $k_{on3}$ and $k'_{off3}$ cannot be independently determined. Simulations performed with $k_{on3}$ values smaller than $10^4$ min$^{-1}$ show accumulation of trimer fractions inconsistent with the experiments. $k_{on3}$ was arbitrarily estimated to $10^5$ min$^{-1}$. The $k_{on3}/k'_{off3}$ ratio was adjusted to account for the measured reaction order. Simulations performed with $k_{on3}$ value larger than $10^5$ min$^{-1}$ displayed the same results when the $k'_{off3}$ rate was proportionally increased. Simulations of the evolution of the TG4T fraction during (TG4T)$_d$ formation in NaCl at 0°C and in KCl at 72°C are displayed in Figures 8 and S4, respectively.

The interpretation of the G-quadruplex formation pathway requires duplex and triplex nucleation centers (D and T) but direct detection and structural investigation
of these species are hopeless due to their extremely weak proportion. Molecular Dynamics (MD) calculation including Na\(^+\) ions performed to investigate the stability of duplex and triplex intermediate species show that the assembly of 2 or 3 dGGGG strands initially aligned as in the (dGGGG\(_2\))\(_4\) canonical quadruplex is unstable and results in ‘cross like’ d(GGGG\(_2\)) and (dGGGG\(_3\))\(_2\) structures that resist to several ns MD simulation (19).

Role of the monovalent ion concentration and temperature on the quadruplex formation rates

In addition to the surface ions that neutralize the backbone phosphorus, \(n-1\) monovalent ions located in the central cavity connect together the successive G quartets of (TG\(_n\)T)\(_4\) quadruplexes (4,20). These monovalent cations have a crucial importance on the quadruplex stability. It seems clear that the preferred potassium coordination in the channel cavity with respect to the other monovalent cations is not directly related to the ions radius but to the lowest dehydration cost of the potassium ion (21,22).

In summary, we find, in agreement with previous investigations (12), that the (TG\(_n\)T)\(_4\) half formation times increase with temperature and when the salt concentration decreases. First, it must be noted that the lifetime of the monovalent cations bound in the G-quartet cavity is extremely short by comparison with the time required for strand association. As shown by Hud et al. (21) the cations bound in the cavity of (G\(_4\)T\(_4\)G\(_4\))\(_2\) exchange with the bulk solvent cations at a rate faster than \(10^{-3}\)s. In line with this result, we observed that addition of 0.1 M KCl to (TG\(_4\)T)\(_4\) in 10 mM NaCl solution results within 3 min in spectral changes reflecting K\(^+\) binding. This clearly establishes that the monovalent cation-binding rate is not a limiting step of the G-quadruplex association pathway. Considering the electrostatic cost of bringing close together negatively charged strands into a multimeric species, it is predictable that the D and T formation rates are slower at low salt concentration. It is also likely that the lifetimes of the D and T intermediates are shorter at high temperature. Hence, high temperature and low salt concentration should reduce the D and T fractions thus resulting in slower quadruplex formation rates. The reduced amplitude at high temperature of the shifts assigned to fast equilibrium exchange between M and D is consistent with this interpretation.

One observes that the formation rates at 60°C of the canonical quadruplexes of (TG\(_4\)T)\(_4\), (TG\(_5\)T)\(_4\) and (rUG\(_4\)U)\(_4\) are comparable within a factor smaller than 10. By contrast, at 0°C in 0.125 M NaCl, the association rates of TG\(_5\)T and UG\(_4\)U are about \(10^8\) times larger than that of TG\(_4\)T (Figures 4 and S1). Fast strand association is systematically observed in the experimental conditions leading to formation of mismatched species. This indicates that the fast strand association rates observed for TG\(_5\)T and UG\(_4\)U at low temperature reflect the faster formation of mismatched structures. The formation of mismatched quadruplexes occurs possibly also in the case of TG\(_3\)T in KCl and of TG\(_4\)T NaCl solution, but due to their shorter lifetime they dissociate rapidly, living unchanged the monomer concentration which vary in that case simultaneously with the canonical quadruplex concentration.

Comparison with previous experimental studies

G-rich strand association into quadruplex structures was until now described by a two-step model: fast equilibrium between monomer and an unstable duplex species followed by duplex association. This model was initially proposed on the basis of the formation-dissociation kinetics of TTG4TT in a buffer containing 137 mM NaCl and 2.7 mM KCl measured by size exclusion chromatography (9). The results of a recent UV study were considered in good agreement with this model (12).

We show in the present study that this model cannot account for the reaction orders determined in the NMR experiments. Qualitatively, there is no marked difference in the monomer association rates and quadruplex dissociation rates determined by UV, chromatography and NMR. The interpretation of the NMR study diverges from that provided by previous studies: (i) about the evaluation of the quadruplex equilibrium constants; (ii) about the interpretation of the fast association rates reported for some oligonucleotides (TG\(_5\)T, rUG\(_4\)U, TG\(_6\)T).

The evaluation of the monomer concentration at equilibrium is not directly accessible to UV measurements. Considering the extremely slow quadruplex dissociation kinetics, it was assumed in the UV study that G-quadruplex formation is ‘kinetically irreversible’ and that the monomer concentration at equilibrium is negligible. Implicitly, the same assumption was done in the chromatography study (9). Accordingly, the reaction order for tetramer formation was derived from the fit of the evolution of the monomer fraction versus time using a model supposing a non-reversible reaction. The reaction order estimated in the chromatography study was 3.8 ± 0.5. In the UV study, the reaction orders were found in the range of 3.4–4.1, in 0.1 M KCl or NaCl solutions.

In the model proposed in the present article, the reaction order for G-quadruplex formation may depend on the monomer concentration [Equation (3)]. The differences in the reaction orders could therefore reflect the difference of oligonucleotide concentration used in the experiments performed by each method. This does not seem the case if one considers that the range of oligonucleotide concentrations examined in the NMR (10–5000 \(\mu\)M), chromatography (100–500 \(\mu\)M) and UV experiments (5–500 \(\mu\)M) are comparable.

Considering the reduced dissociation constants derived from the NMR experiments, it is clear that the free strand concentration at equilibrium cannot be neglected. It follows that the fitting procedure used in the previous studies cannot yield a correct determination of the reaction order. In addition, we consider from our experience, that the fit of the reactant proportion as a function of the time is weakly sensitive to the value of the reaction order. For this reason, we systematically determined the reaction orders by measuring the half reaction
G-quadruplex stability and melting temperature

The slow association and dissociation rates of tetramolecular G-quadruplexes result in marked hysteresis of the UV melting-renaturation profiles making impracticable direct evaluation of the equilibrium melting temperatures (9,12).

The quadruplex equilibrium melting temperatures in 10\(^{-4}\) M oligonucleotide solutions may be determined by two different ways from Figures 4 and S1. As explained above, \(T_m\) in a 10\(^{-4}\) M solution is the temperature for which \(t_{asso,10^{-4}}\) and \(t_{disso}\) are identical. \(T_m\) is also the temperature for which \(F_i = 10^{-4}/2\). The average \(T_m\) values and the difference of the values obtained by each method are listed in Table 1. The equilibrium melting temperatures are at least 20°C lower than the ‘non-equilibrium’ melting temperatures derived from melting experiments performed with a 0.18°C/min gradient (12). This difference is not surprising if one consider the G-quadruplex lifetimes at the melting temperature (2 days to 112 days, Table 1). It is noteworthy that the \(T_m\) computed for (TG4T)\(_4\) in NaCl solution, 32 ± 5°C, is consistent with the value, \(T_m \approx 30°C\), predicted by extrapolation of non-equilibrium melting temperature at infinitely slow temperature gradient (9).

CONCLUSION

The NMR study of the formation of G-quadruplex shows that association of TGnT strands can form multiple tetrameric assemblies. The fully matched canonical quadruplexes are certainly the most stable structures, nevertheless, the faster formation rate of mismatched species and their relatively long lifetimes results in kinetics trapping that may hinder formation of the thermodynamically stable structure during extraordinary long periods.

We have determined the experimental conditions in which the formation of such structures is negligible. The parameters derived from the association–dissociation measurements performed in these conditions show that the canonical quadruplexes are formed via a stepwise pathway involving fast monomer/duplex equilibrium and a triplex intermediate. Triplex formation is limited in 0.125 M KCl solution, due most probably to triplex stabilization by K\(^{+}\). The formation rates of the canonical quadruplexes depend weakly on the nature of the ion and on the G-tract length. On the contrary, the canonical quadruplex lifetimes in K\(^{+}\) solution are 10\(^{6}\) times longer than in Na\(^{+}\) or Li\(^{+}\) solutions and the addition of one G residue in the G-tract increase the lifetime by 4 magnitude orders.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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