Formation of an intramolecular triple-stranded DNA structure monitored by fluorescence of 2-aminopurine or 6-methylisoxanthopterin

Anna K. Shchyolkina, Dmitry N. Kaluzhny, Olga F. Borisova, Mary E. Hawkins1, Robert L. Jernigan1, Thomas M. Jovin2, Donna J. Arndt-Jovin2,* and Victor B. Zhurkin1

Engelhardt Institute of Molecular Biology, Russian Academy of Sciences, 119991 Moscow, Russia, 1National Cancer Institute, NIH, Bethesda, MD, USA and 2Department of Molecular Biology, Max Planck Institute for Biophysical Chemistry, D-37070 Goettingen, Germany

Received June 20, 2003; Revised August 19, 2003; Accepted November 7, 2003

ABSTRACT

The parallel (recombination) ‘R-triplex’ can accommodate any nucleotide sequence with the two identical DNA strands in parallel orientation. We have studied oligonucleotides able to fold back into such a recombination-like structure. We show that the fluorescent base analogs 2-aminopurine (2AP) and 6-methylisoxanthopterin (6MI) can be used as structural probes for monitoring the integrity of the triple-stranded conformation and for deriving the thermodynamic characteristics of these structures. A single adenine or guanine base in the third strand of the triplex-forming and the control oligonucleotides, as well as in the double-stranded (ds) and single-stranded (ss) reference molecules, was substituted with 2AP or 6MI. The 2AP*(T´A) and 6MI*(C´G) triplets were monitored by their fluorescence emission and the thermal denaturation curves were analyzed with a quasi-two-state model. The fluorescence of 2AP introduced into an oligonucleotide sequence unable to form a triplex served as a negative control. We observed a remarkable similarity between the thermodynamic parameters derived from melting of the secondary structures monitored through absorption of all bases at 260 nm or from fluorescence of the single base analog. The similarity suggests that fluorescence of the 2AP and 6MI base analogs may be used to monitor the structural disposition of the third strand. We consider the data in the light of alternative ‘branch migration’ and ‘strand exchange’ structures and discuss why these are less likely than the R-type triplex.

INTRODUCTION

Three-stranded nucleic acid alignments occur in the course of various biological processes, such as recombination or t-loop formation at telomere ends (1). The parallel (recombination) triplex or R-form DNA was predicted theoretically (2) and studied experimentally using protein-free intramolecular (3) and intermolecular (4) oligonucleotide models. Such a triplet hydrogen bond scheme is seen for the triplets G*(C-G) and C*(G-C) in the RNA mini-triple helix formed in the ribosome (5). In principle, the same kind of bonding could accommodate an arbitrary nucleotide sequence with the two identical DNA strands in parallel orientation (2).

The fluorescent base analog 2-aminopurine (2AP) has been used as a site-specific probe of nucleic acid structure and dynamics because it pairs with thymine in a Watson–Crick geometry or makes a mismatch base pair with cytosine (6–12). Upon incorporation of 2AP into double-stranded (ds) B-DNA, its fluorescence is quenched in comparison to the free base (7,13–16). The local conformation and dynamics of mismatches and abasic sites in DNA have been examined using 2AP (9,10,13,14) and it has also been used to study the Mg2+-dependent conformation changes in hammerhead ribozymes (8). Different fluorescence properties of 2AP in the trans A-T base pairs of parallel-stranded DNA and in the cis A-T base pairs of antiparallel-stranded DNA make this base analog a convenient probe for strand orientation in DNA duplexes as well (A.K.Shchyolkina, D.N.Kaluzhny, O.F.Borisova, M.A.Livshits, J.J.Harvey, A.H.A.Clayton, Q.S.Hanley, D.J.Arndt-Jovin and T.M.Jovin, in preparation). 2AP can potentially substitute for adenine in the A*(T´A) triplet, forming a 2AP*(T´A) triplet consistent with the R-form (2,17) (Fig. 1A).

Likewise, the fluorescent guanine analog 6-methylisoxanthopterin (6MI) (18) may mimic guanine in the third strand by forming a 6MI*(C´G) triplet (Fig. 1B) (2,19). Pteridine nucleoside analog probes are highly fluorescent and have been shown to offer another approach to monitor structural perturbation of DNA molecules and subtle DNA interactions with other molecules (20). Here, we demonstrate that the fluorescent base 2AP can be used as a structural probe for monitoring formation of an AT-containing triplex and the fluorescence of 6MI pteridine can be used to probe the structure of a 5’terminal GC-containing triplet.

MATERIALS AND METHODS

Oligonucleotide design

Deoxyribonucleotides potentially able to form the intramolecular R-triplex (RCW, R2APCW, R2APCW-2 and

*To whom correspondence should be addressed. Tel: +49 551 201 1393; Fax: +49 551 201 1467; Email: djovin@gwdg.de
R6MICW) or the duplex (CW and R2APC) were studied (Fig. 2). The GAA linker together with two adjacent GC pairs serve to stabilize the duplex part of RCW, R2APCW, R2APCW-2 and R6MICW, whereas a flexible TTTT loop connects the third strand to the duplex. This design aimed (i) at minimizing displacement of the W strand from the WC duplex by the identical third strand and (ii) at resolving unstacking and dissociation of the third strand from the CW duplex. Deoxyribooligonucleotides NCW and N2APCW served as negative controls for triplex formation, as the nucleotide sequence of the N strand prevents formation of the perfect R-form or of an H and H' type of triplex. In order to test a possible influence of the loop connecting the third strand to the CW duplex part, we studied oligonucleotides R2APCW and R2APCW-2, having -GT4C- and -CT4G- loops, respectively. R2APC and N2APC hairpins and single strands R2AP, R6MI and N2AP were used as controls in the fluorescence experiments for detection of fluorescence signals of 2AP involved in base pairing with thymine and the fluorescent base analogs incorporated into single strands.

In principle, oligonucleotides RCW, R2AP CW, R2APCW-2 and R6MI CW may form not only an R-triplex structure, but also a strand exchange or ‘branch migration’ structure (Fig. 3). These alternatives are considered in Results and Discussion. Oligonucleotides containing 2AP were synthesized and purified by HPLC by the Midland Certified Reagent Co. Inc. (TX). Samples contained 0.7–1 μM oligonucleotide, 0.5 M LiCl, 10 mM Tris–HCl buffer, pH 7.6 ± 0.4 in the temperature range of the melting experiments. 6MI-containing oligonucleotides were synthesized as described (18). Samples were preheated to 90°C and fast cooled on ice. The solution conditions and the method of sample preparation were specifically chosen to ensure the predominant formation of intramolecular structures over intermolecular ones (21).

Fluorescence measurements
Measurement of the fluorescence emission and excitation spectra was carried out using a Photon Technologies International C-60SE spectrofluorometer with automatic correction of the excitation spectra, equipped with a thermostatted cuvette holder.

Thermodynamic analysis of the intramolecular triplex formation
We performed a thermodynamic analysis for the intramolecular binding of the dangling third strand of the triplex-forming oligonucleotide to its double helical CW part. We define such a process as triple helix formation. The transition curve for triplex formation was detected by two experimental techniques. One is the conventional melting curve acquired by measuring the hyperchromicity of the UV absorbance at 260 nm. All nucleotide bases of the third strand absorbing light at 260 nm contribute to this curve, whereas the transition curve measured by 2AP fluorescence emission reflects formation of a single triplet 2AP*(T-A).
The transition curve corresponds to the relationship

\[ A(t) = B(t) + \left[C(t) - B(t)\right] \sigma(t) \]  

where \( t \) is the temperature (°C), \( B(t) \) and \( C(t) \) represent the temperature dependencies of the measured optical signal of the third strand in the bound and the dangling (‘single-stranded’) states, respectively, and \( \sigma(t) \) is shown in equation 2:

\[ \sigma(t) = 1/[1 + \exp[\Delta H/R(1/T - 1/T_m)]] \]  

where \( \Delta H \) is the transition enthalpy, \( T \) is the absolute temperature (\( t + 273.15 \)) and \( T_m \) is the melting temperature (K).

For UV melting curves, \( B(t) \) and \( C(t) \) are defined by \( a(1 - bt) \), the linear dependencies of extinction coefficients of the bases in the bound third strand and in the dangling third strand, respectively.

In the case of the fluorescence melting curve \( B(t) = a \cdot c \cdot (1 - bt) \), where \( c \) is a fraction of fluorescing 2AP in the bound third strand at 0°C. \( C(t) = d \cdot (e^{-m} + m)/(1 + m) \) describes an exponential quenching of 2AP in a dangling third ‘single-strand’ due to collision events with solvent. Parameters \( d, n \) and \( m \) were experimentally determined by ‘melting’ the R2AP single strand, and these values were used during fitting the theoretical curve (equation 1) to the experimental fluorescence melting curve.

Steady-state fluorescence anisotropy measurements and evaluation of relative hydrodynamic volumes of oligonucleotides using EtBr as a probe

To distinguish between intra- and intermolecular three-stranded structures, fluorescence anisotropy measurements were carried out on an SPF-1000 spectrofluorometer (AMINCO), equipped with a thermostatted cuvette holder. The steady-state anisotropy of the fluorescence emission of intercalated EtBr (\( r \)) excited by vertically polarized light was determined from the equation

\[ r = (I_p - I_s)/(I_p + 2I_s) \]  

where \( I_p \) and \( I_s \) are the parallel and perpendicular emission components, respectively. The relative dye concentration did not exceed 1 per 100 nucleotides, thereby avoiding depolarization due to energy transfer. The temperature-dependent contribution of the free EtBr fluorescence to the emission components (equation 3) was taken into account.

The hydrodynamic volume (\( V \)) of molecules with low asymmetry can be estimated by the equation:

\[ V = kT/\tau(1/r - 1/r_0) \]  

where \( \tau \) is the measured fluorescence lifetime of bound EtBr, \( r \) is the measured fluorescence anisotropy and \( r_0 \) is the limiting anisotropy at infinite viscosity (\( T/\eta \to 0 \)), where \( \eta \) is the viscosity of the solution, \( T \) is the absolute temperature and \( k \) is the Boltzmann constant (22). The mean fluorescence lifetimes \( \tau \) of EtBr intercalated in the parallel (recombinant) triplex, conventional antiparallel triplex and calf thymus DNA were found to be practically the same (23,24). This permitted the calculation of a relative hydrodynamic volume (\( V \)) of the oligonucleotides RCW, R2APCW and NCW with respect to the hydrodynamic volume (\( V_{CW} \)) of the CW hairpin (see sequence in Fig. 2):

\[ V/V_{CW} = (1/r_{CW} - 1/r_0)/(1/r - 1/r_0) \]  

for the fluorescence anisotropy values determined under the same solution conditions and temperature (3,25). Under our experimental conditions, one EtBr molecule on average was intercalated in the studied oligonucleotides, causing similar induced structural perturbations. The extension of the double or triple helix on binding one intercalator molecule was
assumed to be 3.4 Å (26,27) and constitutes a uniform systematic error of the estimation of the oligonucleotide hydrodynamic volume using EtBr as probe. An evaluation of the ratio $V/V_{CW}$ offsets the effect of the EtBr-induced extension of the oligonucleotides.

RESULTS AND DISCUSSION

Formation of a triple-stranded structure as monitored by 2-aminopurine fluorescence

The fluorescence emission and excitation spectra of 2AP were found to be different for the triplex-forming (R$^{2AP}$CW), double-helical hairpin-forming (R$^{2AP}$C) and single-stranded (ss) (R$^{2AP}$) oligonucleotides (Fig. 4). As the molar absorption of the three oligonucleotides at the excitation wavelength 310 nm was found to be the same (data not shown), the intensity of the fluorescence emission is proportional to the relative quantum yield (RQY) of 2AP incorporated in the three structures. The excitation maximum for emission at 370 nm for R$^{2AP}$C, R$^{2AP}$CW and R$^{2AP}$ is positioned around 311, 309 and 306 nm, respectively. A blue shift of the excitation maximum reflects an increased exposure to solvent of the 2AP base analog in R$^{2AP}$CW as compared to the ds R$^{2AP}$C (11) and this effect is still more pronounced in the case of the ss R$^{2AP}$. Fluorescence of the 2AP nucleotide is well known to be considerably quenched by base stacking and hydrogen bonding interactions with other bases (compare the R$^{2AP}$ and R$^{2AP}$C temperature profiles in Fig. 5). Thus, by comparing the RQY of 2AP fluorescence as well as the blue shift of the excitation maximum in R$^{2AP}$CW and R$^{2AP}$C (at low temperatures) we conclude that 2AP is more exposed to solvent in the R$^{2AP}$CW structure than in the R$^{2AP}$C duplex. This finding is consistent with relatively weak stacking of bases in the third strand of the ‘R-triplex’ as predicted theoretically, especially for a 5’-pyrimidine-purine-3’ step (see fig. 5B in ref. 2). It should be noted that in the R$^{2AP}$CW construct the 2AP nucleotide is preceded by T, i.e. 2AP is expected to be more accessible to the solvent than in the duplex (Fig. 2). For brevity, we denote

Figure 4. Fluorescence emission and excitation spectra of oligonucleotides at 8°C. R$^{2AP}$CW (filled circles), R$^{2AP}$C (filled triangles) and R$^{2AP}$ (crosses). The excitation wavelength for emission spectra was 310 nm, the emission wavelength for emission spectra was 370 nm. Samples contained 1 µM oligonucleotide, 0.5 M LiCl, 10 mM Tris–HCl buffer, pH 7.6.

Figure 5. Temperature dependence of intensity of the fluorescence emission at 370 nm of 2AP, incorporated in the oligonucleotides. The excitation wavelength was 310 nm. (A) oligonucleotides R$^{2AP}$CW (filled circles), R$^{2AP}$C (filled triangles) and R$^{2AP}$ (crosses); (B) control N$^{2AP}$CW (filled circles), ds N$^{2AP}$C (filled triangles), ss N$^{2AP}$ (crosses); (C) oligonucleotides R$^{2AP}$CW-2 (filled circles) and N$^{2AP}$CW (crosses). Solid curves are the best theoretical fits (see Materials and Methods); dashed curves are the smoothed experimental data. Derived thermodynamic parameters are given in Table 1. Solution conditions as in Figure 4.
the triple-stranded structure stabilized at low temperature as ‘R-triplex’. Whether this is indeed a triplex or rather a branch migration structure (Fig. 3) is discussed in the following two sections, where the experimental data are presented.

The temperature dependencies of the steady-state intensity of the fluorescence emission at 370 nm for the three structures R\(^2\)AP\(_W\), R\(^2\)APC and R\(^2\)AP are shown in Figure 5A. The temperature-dependent quenching of the fluorescence of ss R\(^2\)AP is attributable to collision events with solvent and other species in solution (Fig. 5A, crosses). The emission intensity of R\(^2\)AP\(_W\) displays a sigmoidal curve in the temperature range 8–40°C. Above 50°C, R\(^2\)AP and R\(^2\)APCW display the fluorescence expected for an unpaired 2AP. In the temperature range 8–40°C R\(^2\)AP\(_C\) remains in the double-stranded conformation, with melting starting above 35–38°C (as revealed by the UV temperature profiles; data not shown). The changes in fluorescence of 2AP incorporated into the duplex were minor between 3 and 38°C. The sigmoidal temperature dependence of RQY in the case of oligonucleotide R\(^2\)AP\(_C\) apparently reflects a conformational change in the structure or formation/disruption of the single 2AP\(^*\)(T-A) triplet. This conformational transition occurs in the same temperature range as unstacking of the bases in the third strand, as detected independently by UV hyperchromicity (see below), and therefore can be attributed to melting of the three-stranded structure to a duplex with a ‘dangling’ third strand and unstacked 2AP bases (Fig. 3).

To verify the specific character of this structural transition, we examined the temperature dependence of fluorescence emission of 2AP incorporated in the third strand of the control oligonucleotide N\(^2\)AP\(_C\) (Fig. 5B, filled circles). The third strand N\(^2\)AP of the control is not able to bind specifically to the duplex-forming CW part of the oligonucleotide and does not form runs of base triplets of R- or H-form. Consequently, throughout the temperature range the curves for N\(^2\)AP\(_C\) (Fig. 5B, filled circles) and ss N\(^2\)AP (crosses) coincide. We conclude that 2AP in the third strand of N\(^2\)AP\(_C\) remains in an environment similar to that of 2AP in the single strand. Below 15°C a small decline in the N\(^2\)AP\(_C\) curve as compared to that of ss N\(^2\)AP presumably indicates the presence of non-specific random interactions of 2AP with other bases. The fluorescence of 2AP in ds N\(^2\)AP\(_C\) indicates an involvement in base pairing (Fig. 5B) even though the two strands in this experiment are not fully complementary. The nucleotide sequence of the N\(^2\)AP\(_C\) hairpin allows formation of an 8 bp intramolecular double helix by strand slippage thereby forming a 2AP:T base pair, which explains the temperature dependence of its RQY (Fig. 5B, triangles). Analysis of the N\(^2\)AP\(_C\) UV melting curve (not shown) yielded thermodynamic parameters consistent with this conclusion (Table 1).

Comparison of the data from the triplex-forming oligonucleotide R\(^2\)AP\(_C\) and the triplex-negative control N\(^2\)AP\(_C\) confirms the sequence-specific nature of the fluorescence melting profile of R\(^2\)AP\(_C\) and supports the interpretation of this process as ‘melting’ of the single 2AP\(^*\)(T-A) triplet. The shape of the curve (Fig. 5A, filled circles) agrees with the assumption of a quasi-two-state mode of this transition. Fitting the fluorescence data (see Materials and Methods) yielded the thermodynamic parameters for 2AP\(^*\)(T-A)-triplet formation in R\(^2\)AP\(_C\) tabulated in Table 1.

In general, the structural properties of the nucleotide loop (5’-NTTTTM-3’) that links the third strand to the duplex might affect the thermodynamics or kinetics of melting of the intramolecular ‘triplex’ (28). With this in mind, we compared melting of the two specific triplex-forming oligonucleotides, R\(^2\)AP\(_C\) and R\(^2\)AP\(_C\)-2, containing 5’-GTTTTC-3’ and 5’-CTTTTG-3’ linkers, respectively (Fig. 2). A comparison of the R\(^2\)AP\(_C\)-2 oligonucleotide to the control N\(^2\)AP\(_C\) was also made, as both oligonucleotides have the same 5’-CTTTTG-3’ linker that attaches the third strand to the duplex part (Fig. 5C). The thermodynamic parameters of the R\(^2\)AP\(_C\)-2 triplex are presented in Table 1. In spite of the different G\(^\circ\)C\(^\circ\) and C\(^\circ\)G\(^\circ\)C 3’-end triplets which might influence TT/T loop conformation, we found no difference in either the enthalpy or T\(_m\) of the R\(^2\)AP\(_C\) and R\(^2\)AP\(_C\)-2 triplexes. The data clearly demonstrate that the nucleotide sequence specificity of the third strand and not a difference in the linkers promotes third strand binding to the duplex.

**UV thermal denaturation of the R-forming oligonucleotides**

To correlate the fluorescence signal from the single triplet with the behavior of the third strand, we studied thermal denaturation of the oligonucleotides by UV absorption. Every base of the third strand should contribute to hyperchromicity at 260 nm during ‘triplex’ melting, the contribution of the 2AP analog being smallest as its extinction coefficient at 260 nm is about 0.1 that of adenine (29). Thermal denaturation curves of R\(^2\)AP\(_C\), control N\(^2\)AP\(_C\) and their first derivatives are shown in Figure 6A and B. Denaturation of CW, the ds hairpin, does not contribute to absorption changes below 50°C (Fig. 6A, triangles). The behavior of the R\(^2\)AP and N\(^2\)AP strands in the melting of oligonucleotides R\(^2\)AP\(_C\) and N\(^2\)AP\(_C\) was different (Fig. 6A), differences further accentuated in the first derivatives of these melting curves (Fig. 6B). The R\(^2\)AP\(_C\) melting curve displayed a two-state structural transition, which can be interpreted as melting of a ‘triplex’ to a structure containing the CW hairpin with a dangling third strand, followed by melting of the CW hairpin at high temperature (Fig. 3). In contrast, the first derivative of the N\(^2\)AP\(_C\) melting curve was devoid of a maximum (Fig. 6B, open circles), suggesting a gradual temperature-induced unstacking of the single third strand rather than triplex melting.

To rule out possible intermolecular structures contributing to the observed triplex melting behavior, we measured melting of the R\(^2\)AP\(_C\) oligonucleotide at three different concentrations (0.3, 0.97 and 12 μM). Figure 7 compares thermal denaturation profiles of R\(^2\)AP\(_C\) at 0.97 and 12 μM (Fig. 7A) and at 0.3 and 12 μM (Fig. 7B) detected by UV absorption or 2AP fluorescence, respectively. We found both the UV absorption melting profiles and the 2AP base fluorescence changes to be superimposable at all three concentrations.

Because melting of the CW hairpin continues far beyond the experimental temperature range, treating the R\(^2\)AP\(_C\) melting in terms of a three-state transition from ‘triplex’ to hairpin and then to the open ss oligonucleotide appears impossible (Fig. 6A). One way to extract thermodynamic parameters from the ‘triplex’ melting curve (Fig. 6A, filled circles) is to subtract the duplex CW melting curve (Fig. 6A, triangles). Such an exercise was performed as shown in Figure 6C (filled circles) with the best fit theoretical curve.
enthalpies and melting temperatures are given in Table 1. The formation enthalpy of the `triplex' occurs but rather various duplex con-

Table 1. Thermodynamic parameters of triplex and duplex formation

<table>
<thead>
<tr>
<th>Experimental technique to monitor thermal denaturation</th>
<th>Triplex-forming oligonucleotide</th>
<th>Hairpin-forming oligonucleotide</th>
<th>$\Delta H$ (kJ mol$^{-1}$)</th>
<th>$T_m$ (°C)</th>
</tr>
</thead>
<tbody>
<tr>
<td>2AP fluorescence</td>
<td>$R^{2AP}_{CW}$</td>
<td></td>
<td>$-103 \pm 16$</td>
<td>22 ± 5</td>
</tr>
<tr>
<td></td>
<td>$R^{2AP}_{CW}$-2</td>
<td></td>
<td>$-107 \pm 14$</td>
<td>21 ± 6</td>
</tr>
<tr>
<td>6MI fluorescence</td>
<td>$R^{6MI}_{CW}$</td>
<td></td>
<td>$-76 \pm 10$</td>
<td>17 ± 4</td>
</tr>
<tr>
<td>$A_{260}$</td>
<td>$RC_{CW}$</td>
<td></td>
<td>$-97 \pm 10$</td>
<td>18.0 ± 3</td>
</tr>
<tr>
<td></td>
<td>$R^{2AP}_{CW}$</td>
<td></td>
<td>$-102 \pm 15$</td>
<td>23.5 ± 1</td>
</tr>
<tr>
<td></td>
<td>$R^{6MI}_{CW}$</td>
<td></td>
<td>$-91 \pm 10$</td>
<td>19 ± 4</td>
</tr>
<tr>
<td></td>
<td>$R^{2AP}_{CW}$</td>
<td></td>
<td>$-243 \pm 9$</td>
<td>79.5 ± 0.2</td>
</tr>
<tr>
<td></td>
<td>$N^{2AP}_{CW}$</td>
<td></td>
<td>$-148 \pm 3$</td>
<td>60.6 ± 0.1</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(imperfect 8-bp hairpin)</td>
<td></td>
<td>&gt;85</td>
</tr>
</tbody>
</table>

For the triplex-forming oligonucleotides, the transition from a three-stranded structure to a CW duplex with a dangling third strand was studied. Parameters were determined as the best fits to the theoretical two-state model equation 1 (see Materials and Methods). Experimental conditions: 0.5 M LiCl, 10 mM Tris–HCl buffer, pH 7.6, 20°C.

(equation 1). The parameters from $R^{2AP}_{CW}$ and RCW ‘triplex’ melting are given in Table 1. The formation enthalpies and melting temperatures $T_m$ of the two structures were the same within experimental error, with a tendency for a higher $T_m$ for $R^{2AP}_{CW}$. We obtained remarkably similar parameters for triplex formation derived from the absorption of bases at 260 nm and from the fluorescence of 2AP (Table 1). One may interpret this as a structural quasi-two-state transition with all bases of the third strand simultaneously participating in stabilization of the triple-stranded conformation (in our interpretation, the ‘R-triplex’). The similarity also implies that fluorescence of the 2AP base analog alone may be used to monitor ‘triplex’ integrity.

Formation of a triple-stranded structure as monitored by fluorescence of 6MI pteridine

In the 5'-terminal position of the $R^{6MI}_{CW}$ oligonucleotide, 6MI minimally affects stability of the triple-stranded structure, as revealed by the $T_m$, and apparently allows formation of a 6MI*(C-G) triplet (Fig. 8). Quenching of 6MI fluorescence is known to take place upon stacking with other bases (18) and proceeds upon cooling in the case of $R^{6MI}_{CW}$ but not in the case of ss $R^{6MI}$ (Fig. 8). The formation enthalphy of the ‘triplex’ derived from the melting curve in Figure 8 (filled circles) was ~25% less than that for RCW and $R^{2AP}_{CW}$ (Table 1). This may be accounted for by a terminal position of the triplet under consideration. Furthermore, the electronic properties of 6MI are markedly different from those of guanine, which may alter the energy contribution of stacking interactions of 6MI*(C-W) with the adjacent triplet. Moreover, minor structural perturbations of the triplet terminus due to substitution of 6MI for G could weaken ‘triplex’ stability. Nevertheless, formation of a terminal 6MI*(C-G) triplet as measured by fluorescence of this base analog is an additional argument in favor of the R-like triplex structure.

Alternative triple-stranded structures

Figure 3 shows some alternative structures possible where no triplex occurs but rather various duplex configurations are formed by ‘branch migration’ or ‘strand exchange’. Our biphasic melting data, in which the higher temperature melt is not affected by the presence of a third strand (see Table 1 and Fig. 6), is not compatible with various duplex structures whereby the R strand invasion would shorten the 12 bp CW duplex. The latter should lead to a broader and lower temperature melting curve. Furthermore, we demonstrated a direct involvement of the terminal base in stabilization of the triple-stranded structure by using 6MI substitution, which strongly undermines a model with fraying ends.

If the structure is rather an equilibrium between two ‘strand exchange’ duplexes (Fig. 3), the first step of the melting curve should be assigned to melting of the shorter 10 bp RC hairpin and the second step to melting of the 12 bp CW hairpin (as described above). As we demonstrated, an RC hairpin unfolds closer to 80°C than to the low temperature transition of ~20°C (Fig. 6).

Neither the branch migration nor the strand exchange structures alone are consistent with the experimental data. On the other hand, we cannot exclude that an equilibrium may exist at low temperature between the R-triplex and the other structures. Nevertheless, estimation of the hydrodynamic volume of the RCW and $R^{2AP}_{CW}$ oligonucleotides (see below) suggests close alignment of the third strand along the CW duplex at this low temperature such that no true ‘dangling strand’ can be present.

Estimation of the hydrodynamic volume of the oligonucleotides by intercalated dye fluorescence anisotropy

The fluorescence anisotropy of an intercalated DNA dye reflects the hydrodynamic volume of the DNA and, thus, is a probe for the intra- or intermolecular nature of the structures. The steady-state fluorescence of EtBr intercalated in short DNA duplexes (21,30), triplexes (3) and tetraplexes (31) has proved to be a convenient tool for the detection of multistranded structures (see Materials and Methods). EtBr does not destabilize DNA triplexes at low occupancy (23). The effect of EtBr intercalation on the melting curves of $R^{2AP}_{CW}$ is shown in Figure 9A. As expected, binding of one EtBr molecule (on average) to $R^{2AP}_{CW}$ had a minor effect on its stability and did not influence the value of the absorption hyperchromicity. The measured steady-state fluorescence anisotropy values were used to calculate ratios of the hydrodynamic volume of RCW or $R^{2AP}_{CW}$ to that of CW at different temperatures according to equation 5 and are plotted in Figure 9B. The resulting ratios ($V/V_{CW}$) at 6°C are 1.4–1.5. These values are very close to

\[ V/V_{CW} = \frac{V_{RCW}}{V_{CW}} = \frac{V_{R^{2AP}_{CW}}}{V_{CW}} \]
those found in an earlier study (32) for the ‘canonical’
antiparallel purine intramolecular triplex anti-AG [5’-d(CT)5-
L-d(AG)5-L-d(GA)5-3’] of $V/V_{CW} = 1.40$ and for the parallel
‘recombination-like’ triplex par-AG [5’-d(GA)5-L-d(TC)5-L-
d(GA)5-3’] of $V/V_{CW} = 1.45$, where L stands for the linker
pO(CH$_2$CH$_2$O)$_3$p. The ratio $V/V_{CW}$ has been shown to be
sensitive to formation of intermolecular structures, for
example in the presence of magnesium ions (33). In such a
case, dimerization of the anti-AG or par-AG oligonucleotides
(mentioned above), increases the value of $V/V_{CW}$ up to 2.0–2.3
(33). Thus, we interpret the ratios for the data at 6
°C (Fig. 9B) as reflecting an intramolecular triple-stranded structure with
the third strand tightly arranged along the double-helical part
of the oligonucleotide. Importantly, the similarity of the
thermal denaturation curves of R2APCW at concentrations of
0.3, 0.97 and 12
$m$M independently supports the assumption
that the structures are intramolecular in nature (see Fig. 7).

The $V/V_{CW}$ ratios for R$_2$APCW and RCW increased with
temperature to a value of 1.60–1.64 at ~26°C, after which the

Figure 6. Thermal denaturation of R$_2$APCW and control oligonucleotide
N$_2$APCW monitored by absorption at 260 nm. (A) Melting curves of R$_2$APCW
(filled circles) and of N$_2$APCW (open circles), left ordinate; melting curve of
CW hairpin (filled triangles), right ordinate; (B) first derivatives of the
melting curves of R$_2$APCW (filled circles) and N$_2$APCW (open circles);
(C) determination of the thermodynamic parameters of R$_2$APCW triplex
formation from the absorption thermal denaturation experiments. The triplex
melting curve (filled circles) is a result of subtraction of the CW contribution
to the melting profile of R$_2$APCW (see filled circle and filled triangle in curves
Fig. 5A). The solid curve is the best theoretical fit for a two-state model (see
Materials and Methods). The derived thermodynamic parameters are given in
Table 1. The concentration of each oligonucleotide was 0.77 
$m$M.

Figure 7. Thermal denaturation of R$_2$APCW at different oligonucleotide
concentrations. (A) UV absorption melting profiles. R$_2$APCW concentrations
were 0.97 (filled circles) and 12 
$m$M (filled triangles). The 12 
$m$M sample
was measured in a 1 mm cell and the plotted OD values are calculated for a
1 cm path. (B) Temperature dependence of 2AP fluorescence in R$_2$APCW at
370 nm: oligonucleotide concentrations were 0.3 (filled circles) and 12 
$m$M
(filled triangles).
value decreased to 1.05±1.1 at temperatures of 48–50°C (Fig. 8B). We interpret the data as follows. Below 26±28°C the third strand of the RCW and R2APCW oligonucleotides remains attached to the duplex, presumably due to a triplet-stabilizing effect of the EtBr, though local bulging of bases may cause the slight increase in $V/V_{CW}$ observed. At temperatures >30°C the triplex third strand begins to melt, as shown in Figures 5 and 6, decreasing contact with the duplex. Correspondingly, the EtBr molecules would redistribute from triplex binding sites to duplex sites. Above 48–50°C we would expect the third strand to be freely rotating relative to its duplex part, thereby contributing little to the hydrodynamic volume of the complex and thus the anisotropy value diminishes to a value similar to that for the CW duplex. These data further support the intramolecular nature of RCW and R2APCW triplex formation at low temperatures and the melting observed by spectroscopic means presented in the previous sections.

Concluding remarks

Compared to conventional antiparallel triplexes Py*(Pu´Py), the parallel three-stranded structure [or putative recombination R-triplex (2)] is less stable (3), therefore its experimental detection is more complicated. The structure involving the R strand is relatively weak, with a formation enthalpy of −100 kJ (i.e. ~2.5-fold less than the enthalpy of the RC duplex; see Table 1). In other words, the low temperature transition is principally different from the duplex melting. The low stability of the ‘R-triplex’ is probably related to less favorable base–base interactions (compared to Hoogsteen-type interactions) with heterogeneous and weak stacking in the third strand. In addition to the planar base–base interactions, the stereochemistry of R-triplets allows for formation of inter-planar interactions between the third strand bases and the Watson–Crick duplex (2). Such a dynamic behavior and relatively low stability of the base triplets, however, are major factors making the ‘R-triplex’ an ideal candidate for a transient intermediate in homologous recombination (2,34–36) or in t-loop formation at the mammalian telomere ends (1). In this study we have used two different fluorescent probes that enabled us for the first time to follow the behavior of the individual R strand in the course of triplex melting. Importantly, the oligonucleotides RCW and R2APCW were specially designed to have exceptionally stable CW hairpins, which facilitated equilibrium formation of the R-type triplex at low temperature. Although we cannot completely exclude formation of alternative (non-R-triplex) structures, our data
strongly disfavor ‘branch migration’ or ‘strand exchange’ structures under our experimental conditions. The results demonstrate the potential of 2AP and 6MI as structural probes to monitor the rearrangement of nucleic acid strands upon formation and dissociation of multi-stranded complexes.

ACKNOWLEDGEMENTS

The authors appreciate the helpful discussions of V. L. Florent’ev, M. Livshits, I. Panyutin and N. Ulyanov and the assistance of D. Cherny with the figures. The study was supported by RFBR grants nos 01-04-48561 and 02-04-49182, a Collaborative Linkage NATO grant SA (LST.CLG.978296) and the Max Planck Society.

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