

The Effect of a G:T Mismatch on the Dynamics of DNA

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Abstract

Distortions in the DNA sequence such as damages or mismatches are specifically recognized and processed by DNA repair enzymes. A particular challenge for the enzymatic specificity is the recognition of a wrongly-placed native nucleotide such as thymine in T:G mismatches. An important step of substrate binding which is observed in many repair proteins is the flipping of the target base out of the DNA helix into the enzyme's active site. In this work we investigate how much the intrinsic dynamics of mismatched DNA is changed compared to canonical DNA. Our molecular dynamics simulations of DNA with and without T:G mismatches show significant differences in the conformation of paired and mismatched DNA. The wobble pair T:G shows local distortions such as twist, shear and stretch which deviate from canonical B form values. Moreover, the T:G mismatch is found to be kinetically less stable, exhibiting two states with respect to base opening: a closed state comparable to the canonical base pairs, and a more open state, indicating a proneness for base flip. In addition, we observe that the thymine base in a T:G mismatch is significantly more probable to be flipped than thymine in a T:A pair or cytosine in a C:G pair. Such local deformations and in particular the existence of a second, more-open state can be speculated to help the target-site recognition by repair enzymes.

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Introduction

Deamination of cytosine or methyl-cytosine is a DNA damage resulting in mutation to uracil or thymine, respectively, thus leading to G:U or T:G mismatches. Glycosylases such as the human thymine DNA glycosylase (TDG) or uracil DNA glycosylase (UDG) recognize T:G or G:U mismatches and remove specifically the mismatched T or U, respectively. Detection of a T:G mismatch means recognition of one of the four nucleotides which compose the DNA but is positioned at a wrong site, hence resulting in sequence mismatches.

From the many structures of glycosylases complexed to damaged DNA [1,2] it is known that damaged, mismatched or wrong bases are flipped out of the helical DNA duplex into the enzyme's active site. This base flip has been argued [3,4] to facilitate access to proton acceptor groups of the scissile base, i.e. the base which will be removed by the repair enzyme. In the debate how glycosylase enzymes recognize a damaged or mismatched base two mechanisms are discussed. One is a "passive" mechanism in which the enzyme detects extra-helically exposed, already (at least partially) flipped-out bases. This mechanism implies that base pair opening up to several degrees of flipping is more likely for damaged/mismatched bases than for intact canonical ones. The alternative mechanism involves flipping of the base while the enzyme travels along the DNA, relying on the enzyme specifically enhancing the flip-out of its target base [4,5].

Solution NMR studies have shown that the T:G mismatch introduces only local perturbations to the DNA B form not extending beyond the neighbouring base pairs [6]. Besides small deviations in the backbone torsion angles the authors report an

asymmetry of the λ -angles between the glycosidic bonds and the base pair vector (C1'-C1') for T:G mismatches as opposed to the rather symmetric λ angles observed in canonical base pairs [6].

An experimental probe for base opening and re-closing of hydrogen bonded base pairs is the exchange of imino protons of guanine, uracil or thymine which can be measured by NMR [7–9]. The base-opening rate can be calculated from the imino proton exchange rate assuming that both rates are equal if the exchange itself is fast (which can be achieved by the use of proton accepting catalysts). However, imino proton exchange rates cannot be directly used as a measure for base flipping since solvent accessibility of the imino protons can be achieved already at low flip (opening) angles [10–12]. Moreover, as imino-proton accessibility can be achieved by flipping of either of the two bases in a pair, the kinetics of a single base flipping completely out of the DNA double helix - towards the conformation which has been observed in complexes with DNA repair proteins - can thus not be directly extracted from the experiments.

Molecular simulations have proven to be a powerful tool for obtaining information on structure and dynamics at the atomic level which is not directly accessible to experiments, and have been used successfully to analyse conformational changes in proteins and DNA [13–27].

Sequence dependent dynamics of uncomplexed DNA with different types of damage, lesion or mismatches, has been investigated by numerous molecular dynamics studies [15,28–37]. Simulations of base flip have been successfully conducted on free DNA [28–40] and in complex with different DNA repair enzymes [21,22,41,42].

Crystallisation experiments on human Thymine DNA glycosylase solved the structure of a protein-DNA complex with a flipped-out product analogue [43]. Kinetic experiments showed the importance of conserved residues for base flipping [44]. Recent work by the same group revealed the structure of a DNA with a substrate-analogue (U^F) [45] complexed to wild-type and mutant protein. The authors moreover report MD simulations of the protein-DNA complex in the flipped-out form, showing that two conserved residues destabilise the completely-flipped form of target dT as opposed to substrate dU. They conclude this incomplete flip to reduce the accessibility of the dT, which could minimise aberrant T removal from A:T pairs. This is consistent with earlier biochemical work by the same authors [46] in which the reversible nucleotide flipping was found to be much more rapid for G:T than for G:U substrates.

Since flipping of a DNA base occurs on time scales much larger than accessible by direct MD simulations various flavours of enhanced molecular dynamics have been used which all force the base flip to occur by applying an external potential. The main difference is the definition of the reaction coordinate, i.e. the geometric parameter (e.g. internal coordinate) being restrained. For a detailed review of different methods see [29,40]. Comparisons of the popular force fields has been reported in [33]. The two popular force fields CHARMM [47] and AMBER parm99 [48] show similarly good agreement with experimental imino proton exchange data. However, the detailed atomic picture as well as the calculated PMFs vary between the different force fields. In a more recent study [35] long simulations of long DNA duplexes with the CHARMM force field and with the improved Amber force field parmbsc0 [49] have been compared. The authors conclude a “very remarkable similarity between parmbsc0 and CHARMM27 estimates” for helical stiffness. Hydrogen bonds between Watson-Crick pairs C:G and T:A are found to be less strong with CHARMM27 compared to parmbsc0. Transient loss of hydrogen bonds is observed to be common using either of the two force fields. In conclusion both force fields are of similar quality to obtain a “consensus picture of the basic structural dynamics characteristics of B-DNA” [35].

The opening of T:G pairs in DNA (and G:U in RNA) has been studied recently, by performing a combination of imino proton exchange measurements and molecular dynamics simulations [12]. The authors defined opening by a linear combination of the flip angles of the two bases G and T. The energetically most favourable opening pathway was found to be a coupled rotation of both bases, opening through the major groove. Moreover, the authors conclude that the common two-state model for base pair opening can be applied since imino protons of the closed pair are found to be not accessible for the solvent. However, proton exchange was reported to take place with only 10–40% accessibility [12].

In this work, we apply molecular simulations to explore how much the intrinsic dynamics of the mispaired DNA, compared to intact, well-paired DNA, contribute to mispair recognition by repair proteins. We have performed molecular dynamics simulation of short sequences of unbound DNA in water in order to investigate the dynamics of the paired DNA and of DNA carrying one T:G mispair instead of a G:C pair or an A:T pair. We have analysed the conformational difference of the DNA at the T:G mispair compared to Watson-Crick pairs C:G and T:A. Furthermore, we have computed the free energy for the non-enzymatic flip process in water in order to investigate whether a thymine from a T:G mispair can be flipped out of the DNA double helix more easily than flipping cytosine from C:G or thymine from T:A.

Methods

System Setup

Three setups of DNA oligonucleotides of 17 base pairs length were prepared in standard B-DNA form, d(GCTCTGTACGT-GAGCAG), the site of interest is underlined. This is the part of the DNA sequence observed in the crystal structure of the hTDG-DNA complex (2RBA [43]) where the protein is bound to. The site which is abasic in the hTDG-DNA complex has been modelled with either cytosine (C:G) or thymine (T:G), both flipped in. As another reference, a third setup with an T:A pair at the C/T:G site was prepared. These initial models were build and minimised with CHARMM [47]. The CHARMM 27 Force field [50] was used throughout.

The 17-bps oligomers have a length of ~ 60 Å and a width of ~ 20 Å. The systems were solvated with explicit water, using the TIP3P model [51], extending to at least 10 Å beyond the DNA in each direction in a cubic box ($x = y = z = 90$ Å). The cubic shape ensures that even after rotation there would be enough distance between two adjacent images. 36 Na^+ counter-ions were added to neutralize the system and an excess of Na^+ and Cl^- ions to obtain a physiological concentration of 150 mM NaCl. The addition of the ions was carried out by random substitution of water oxygen atoms.

Simulations were performed using periodic boundary conditions and the long-range electrostatic interactions were treated using the Particle Mesh Ewald method [52] on a $92 \times 92 \times 92$ charge grid, with a non-bonded cutoff of 12 Å. The short range electrostatics and van der Waals interactions were truncated at 12 Å using a switch function starting at 10 Å.

The solvated structures were minimized using 5000 steps of steepest descent, followed by minimization with the conjugate gradient algorithm, with solute atoms harmonically restrained until an energy gradient of 0.01 kcal/(mol Å) was reached. The system was then gradually heated for 30 ps to 300 K with 1 K temperature steps with harmonic restraints on the solute atoms.

The systems were equilibrated in three different stages with the numbers of particles, pressure (1 bar) and temperature kept constant (NPT ensemble) during 75 ps. In the first 25 ps velocities were rescaled every 0.1 ps and in the second 25 ps Langevin dynamics were used to maintain constant temperature. Pressure control was introduced in the third 25 ps and in the production run using the Nosé-Hoover Langevin piston with a decay period of 500 fs [53]. The harmonic restraints were gradually lifted (to 0.5, 0.25 and 0.05 kcal/(mol Å²)) in the three equilibration stages.

Unbiased MD Simulations

After equilibration, unbiased NPT production runs were performed for 60 ns. The integration time step was 2 fs and coordinates were saved with a sampling interval of 2 ps. All covalent bonds lengths involving hydrogen atoms were fixed using SHAKE algorithm [54].

Several independent MD simulations were carried out by assigning different initial distributions of starting velocities to the minimized systems: three runs for the two setups of paired DNA (C:G and A:T), and five for the mispaired T:G model (cf. Table 1).

Biased (ABF) MD Simulations

For the simulation of the base flip we applied the Adaptive Biasing Force (ABF) method [55–57]. In ABF the reaction coordinate is discretized into small bins. Sampling is carried out along the reaction coordinate in a continuous fashion. In each bin samples of the instantaneous force acting along the reaction coordinate are accrued up to a certain threshold. If this threshold

Table 1. List of MD simulations.

	unbiased	ABF
C:G	3*60 ns	3*30ns
T:G	5*60 ns	5*32ns
T:A	3*60 ns	3*30ns

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is reached the adaptive biasing force is applied so as to “drive” the system into the next bin. The reaction coordinate for the base flip has been defined as a pseudo-dihedral angle between the flipping base, the sugar moiety of the same nucleotide, the sugar of the next nucleotide, and the base of the next nucleotide plus the base and sugar of the opposing nucleotide downstream (see Figure 1). This definition of the flipping coordinate is similar to the one proposed and applied in [31,33,38]. The potential of mean force (free energy profile) was obtained by discretising the reaction coordinate between 10° and 180° into windows of 2° width, and in each window 2000 samples were collected before the bias was applied. For C:G and T:A we carried out three ABF simulations each, and for T:G five individual ABF simulations were started with different initial velocities. The biased simulations were run for 24 ns each (cf. Table 1). All molecular dynamics simulations were performed with the NAMD [58] program.

Analysis

To calculate potentials of mean force, angles were binned by 2 degree and translation parameters were binned by 0.2 Å. The free energy difference, ΔG , to the reference state was evaluated according to

$$\Delta G = -RT \ln \frac{P(i)}{P(ref)}$$

where $P(i)$ is the probability of finding the system in state i and $P(ref)$ is the probability of finding the system in the reference state. Probabilities have been calculated from the number of occurrences within a bin. The bin with the highest occurrences has been chosen as the reference state. The free energy has been evaluated in the 99% confidence interval.

For all analyses (unbiased and ABF simulations), properties were evaluated for each run individually. Then the averages and standard errors over the respective individual runs were calculated.

In the analyses of the unbiased MD simulations, the first 10 ns of each trajectory were not included. The Root mean Square Deviation (RMSD) as a function of time, plotted in Figure S1 in the supplementary material suggests this simulation time to be sufficient. The convergence of these simulations was furthermore evaluated by comparison of the properties computed from different simulation lengths, i.e. 40, 50 and 60 ns (shown as Figures S5–S17 in the supplementary material). Convergence of the ABF simulations has been evaluated in a similar manner by

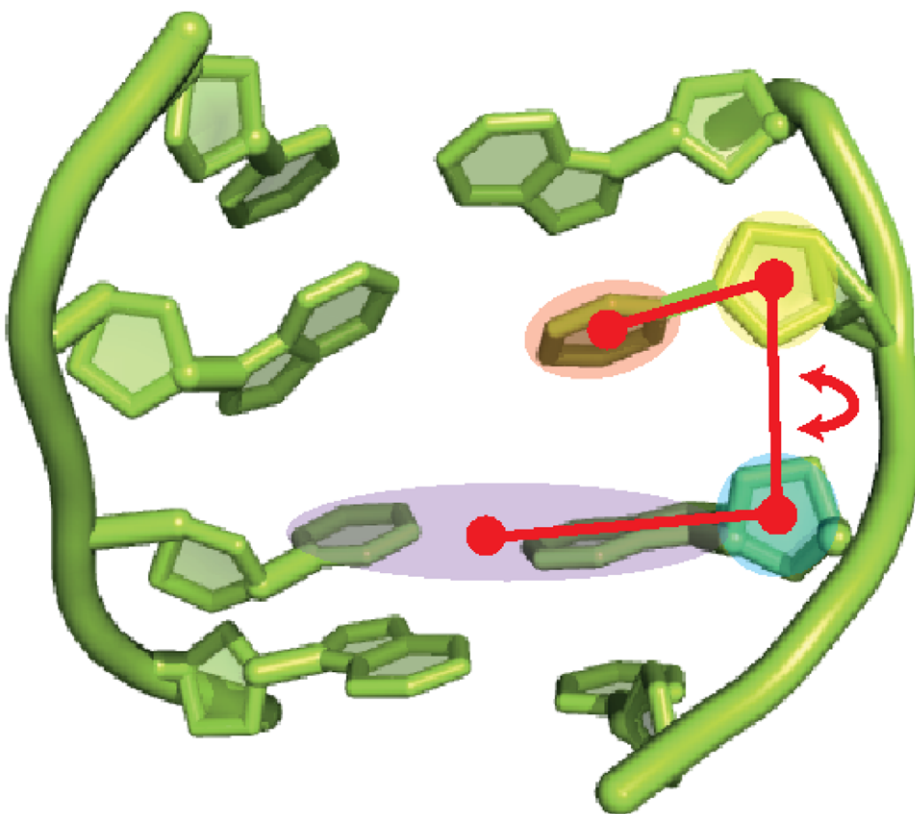


Figure 1. Definition of the reaction coordinate for the base flip simulations: the flip angle is a pseudo-dihedral between the centres of mass of the flipping base (red shade), the sugar moiety of the same nucleotide (green shade), the sugar moiety of the next nucleotide (yellow shade), and the base of the next nucleotide plus the complementary base in the other DNA strand (blue shade). doi:10.1371/journal.pone.0053305.g001

comparing the free energy profiles obtained after 20 ns–32 ns simulation time (cf. Figure S19 in the supplementary material).

The conformations of the paired and mispaired DNA were characterized by calculating twelve helical parameters, six for base steps (the three rotational parameters: roll, tilt and twist, and the three translational parameters: slide, rise and shift) and another six for base pairs (buckle, propeller, opening, stagger, shear, stretch) that define the local DNA geometry. In addition, we have computed the lambda angles which define the angles between the glycosidic bonds (N1/N9–C1') and the base-base vector (C1'–C1').

Hydrogen-bond occupancies were calculated as the ratio of the time when the hydrogen bond is formed to the total time of the trajectory. Two atoms are considered here to form a hydrogen bond if the acceptor-donor distance is <3.0 Å and the acceptor-hydrogen-donor angle is $>135^\circ$.

Solvent accessible surface areas have been computed by placing a probe sphere of radius $r_{vdW}+1.4$ Å in contact with the atomic van der Waals sphere, both centred at the atom. The parts of the surface spheres where the centre of the spherical probe can be placed without penetrating other atoms add up to the solvent accessible surface area [59].

The Molecular Dynamics simulations have been carried out using the program NAMD2.7 and applying the CHARMM27 force field. All simulations have been performed on the local group cluster, the Heidelberg Linux Cluster (HELICS) and the North-German Supercomputing Alliance (HLRN).

All molecular images were generated with the molecular visualization program VMD [60] and with the molecular graphics program Pymol [61]. Structural analysis was performed using standard programs; Curves5.3, gromacs-4.5.5 [62–64] tools and our own scripts.

Results

DNA Conformation

We have examined the conformation of the DNA double helix carrying the mispair and compared it to the intact DNA, analysing the local conformation at the mispair T:G, the pair C:G and for comparison at a T:A pair (see Figure 2 for a schematic drawing of the three base pairs). Figure 3, and Figures S1 and S2 show the free energy profiles for the local helical parameters representing the local DNA conformation at the T:G, C:G and T:A base pair, respectively. Among the helical parameters characterizing the base step only the twist angle and the shift translation exhibit significant differences between the C:G or T:A pairs and the mispaired T:G. The twist angle has a free energy minimum at $32 \pm 1^\circ$ and $30 \pm 1^\circ$ for the two Watson-Crick pairs, C:G or T:A, respectively, whereas for the T:G wobble pair a higher twist angle ($39 \pm 1^\circ$) is more probable. In case of the shift translation, the T:G mispair shows two free energy minima. The first one is located at around -0.5 ± 0.2 Å, at about the same position as the free energy minimum of the C:G shift (-0.4 ± 0.1 Å) and close to the 0.2 ± 0.1 Å shift of the T:A pair. The second free energy minimum of the T:G mispair, which is only marginally higher in energy than the first one, is observed at -2.3 ± 0.4 Å.

The free energy profiles of the base-pair parameters computed from the unbiased simulations of the three different DNA setups again show high similarities between the two Watson-Crick pairs C:G and T:A (see Figures 3, S2 and S3). The most pronounced differences between Watson-Crick pairs and the wobble pair can be observed for the shear and stretch translations, and for the opening angle. T:G shows a free energy minimum for shear at -2.3 ± 0.1 Å and the most probable stretch translation at -0.5 ± 0.05 Å, which deviate from the value of the Watson-Crick

pairs by -2 Å and -0.5 Å, respectively. Moreover, in the T:G case, a second free energy minimum for the base-pair stretch (0.4 ± 0.1 Å) can be observed, albeit with higher statistical errors. The free energy profile of the base pair opening angle also shows that the T:G wobble pair has (at least) two states. The most likely state has an opening angle similar to that of the Watson-Crick pairs (-1.0 ± 0.5 Å). However, a second, slightly less probable state which is separated by only 1.5 kcal barrier, is observed at an opening angle of $45 \pm 1^\circ$ (Figure 3). The distortions on the DNA carrying the T:G mispair are very local as can be seen from the comparison of the local parameters of the flanking bases and base pairs (supplementary material, Figures S5, S6, S7, S8, S9, S10, S11, S12, S13, S14, S15, S16). The only parameter of neighbour which is affected is the shift of the base step preceding the mispaired T. This is due to the definition of the shift, i.e. the displacement along the x-axis with respect to the neighbouring base step. The displacement of base number 10 (T) leads to a shift with respect to base number 11 and base number 9.

Hydrogen Bonds

Table 2 lists the occupancies of hydrogen bonds of the target base with the base on the complementary DNA strand or with solvent water, respectively. As anticipated the hydrogen bonds in the Watson-Crick pairs C:G and T:A are very stable as manifested by the hydrogen bond occupancies of $78 \pm 1\%$ and $96 \pm 1\%$. Only the C-N4–G-O6 bond in the C:G pair is more dynamic and is formed $68 \pm 5\%$ of the simulation time. The T:G mispair shows one very stable hydrogen bond between G-N1 and T-O2 which is formed $72 \pm 3\%$ of the simulation time. Another hydrogen bond to T-O2 is formed by G-N2 with an occupancy of $34 \pm 6\%$, suggesting that the T-O2 fluctuates between the two hydrogen bonded states (see Figure 4).

T-N3 is observed to form hydrogen bonds to G-O6 about half of the simulation time ($50 \pm 6\%$ occupancy). The imino proton (N3-H) of thymine also forms hydrogen bonds to solvent water, for about the same percentage of the simulation time as the T-O2–G-N2 hydrogen bond is occupied ($35 \pm 7\%$ and $34 \pm 6\%$, respectively).

The analysis of the hydrogen bonds with water shows that all oxygen atoms involved in hydrogen bonds within the base pair (C-O2, T-O2 and T-O4) accept additional hydrogen bonds from solvent water. The amino group nitrogen atom C-N4 acts as hydrogen bond donor to solvent water, too. A significant difference, however, is the observation of hydrogen bonds between T-N3 and water in the mispaired Thymine only.

Base Flip

Figure 5 shows the free energy profile of the pseudo dihedral flip angle for the three setups T:A, T:G, and C:G computed from the unbiased MD simulation. Whereas the Watson-Crick base pairs C:G and T:A exhibit a rather narrow free energy minimum around $37 \pm 1^\circ$ and $38 \pm 1^\circ$, two free energy minima can be observed in the free energy profile of the mispaired T:G DNA. The most probable flip angle is at $47 \pm 1^\circ$ and a second free energy minimum, about 0.3 kcal/mol higher in energy is located at $68 \pm 1^\circ$. The rather low barrier between the two free energy minima allows for frequent transitions to be observed (cf. Figure S18).

By applying the technique adaptive biasing force, we have computed the free energy for the rotation (flip) of a single base out of the DNA double helix up to 180 degree flip angle. A complete rotation of the DNA base, including passage of the minor groove turned out to require too high forces resulting in deformation of the DNA. Therefore, we have computed the potential of mean

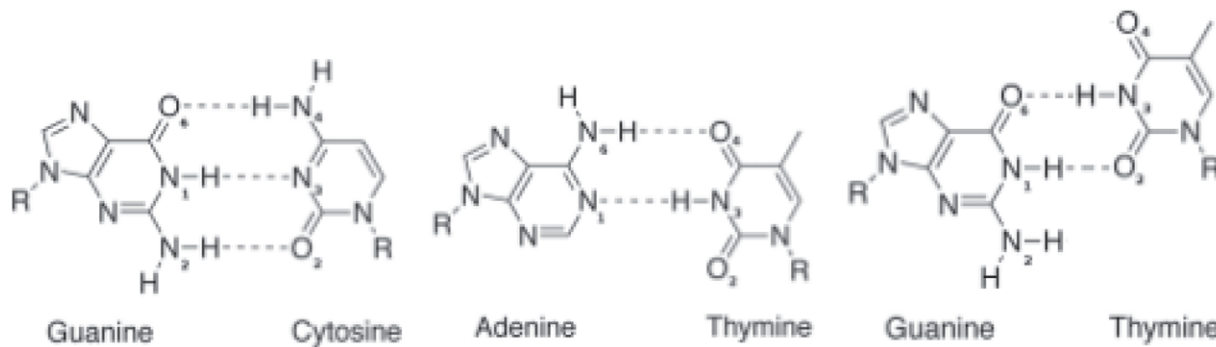


Figure 2. Schematic drawing of Watson-Crick pairs C:G (left) and T:A (middle), and the mispair T:G (right).
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force only for a flip through the major groove. The free energy profile is plotted in Figure 6.

The positions of the free energy minima are the same as those observed in the unbiased simulation for the two Watson-Crick base pairs. The thymine base of the mispaired T:G, however, exhibits two free energy minima at $42 \pm 6^\circ$ and at $67 \pm 2^\circ$, separated by a barrier of ~ 1 kcal/mol. These free energy profiles are comparable to those obtained from the unbiased simulation (Figure 5).

As anticipated, the flip of a cytosine base from a C:G pair requires significantly more energy (on average 11 ± 2 kcal/mol) than the flip of a mispaired thymine (5.5 ± 1.8 kcal/mol). The flip of a thymine base from a T:A pair has only slightly lower free energy barrier but a significantly narrower error range (10.4 ± 0.5 kcal/mol) than the cytosine flip of the C:G pair.

Water Accessibility

In order to analyse how the base flip affects the otherwise shielded intra base-pair hydrogen bonds we have computed the water accessibility of the hydrogen bond formed with atoms of cytosine and thymine (N3, O2, N4, and O4, respectively) as a function of the base flip angle. The respective curves are plotted in Figure 7, left. The additionally computed water accessible surface area of those atoms is shown in Figure 7, right. Both, hydrogen bonds and solvent accessible surface area show a similar dependency on the base flip angle, except for the O4/N4H2-atom. For the N3-atom both curves show minima at about the same flip angle ($35 \pm 5^\circ$ for C:G and T:A, and $40 \pm 5^\circ$ for T:G, respectively) as the free energy profile. However, the minima of hydrogen bonds with the N3 atom and solvent accessible surface area of the T:G mispair are significantly narrower than the free energy minimum of the flip angle. T:G exhibits an average number of 0.3 hydrogen bonds between the N3-atom and solvent water even at the free energy minimum flip angle ($40 \pm 5^\circ$) which is in agreement with the $35 \pm 7\%$ hydrogen bonds occupancy observed in the unbiased simulations (cf. Table 2).

In the two cases where O2 is hydrogen-bonded to guanine (G:C and T:G) the water-O2 interaction (either in terms of numbers of hydrogen bonds or as solvent accessible surface area) is minimal at a flip angle of $70 \pm 10^\circ$ and $60 \pm 10^\circ$, respectively. The O2-atom of thymine is not involved in a Watson-Crick interaction when paired to adenine, and is then able to form hydrogen bonds with solvent water in a flipped-in state. Hence, the number of hydrogen bonds does not change significantly with respect to the flip angle. Its solvent accessibility, however, is minimal at the flipped-in state ($45 \pm 10^\circ$ flip angle) and increases with flip angle similar to the free energy for base flipping.

Atoms O4 and N4 show an increase of numbers of hydrogen bonds and an increase of the solvent accessible surface area at a flip angle of up to $60\text{--}70^\circ$. The exceptions are the number of hydrogen bonds formed between the mispaired thymine O4 and solvent water, and between N4H2 with water via the second hydrogen atom of, which both are already present in the flipped-in state (see also Table 2).

Figure S21 in the supplementary material shows snapshots of the base flip trajectories. In the flipped-in state the thymine/cytosine base forms two/three hydrogen bonds to its complementary base. At a flip angle of $60\text{--}70^\circ$ only the O2-atom is buried in the DNA double helix and forms a hydrogen bond to the opposite base on the complementary strand. At 180° flip angle, the base is completely flipped out into the solvent, whereas neighbouring bases are properly paired.

Discussion

The analysis of the DNA helical parameters clearly shows a distortion in the DNA containing the T:G mispair compared to canonical DNA. In particular the base pair parameters shear and stretch show a significant deviation from the values observed for the Watson-Crick pairs. A higher twist angle is another indicator for the mispaired base. The wobbling of the T:G mispair is manifested in several respects. The hydrogen-bond occupancies indicate that the O2-atom of T alternates between being hydrogen bonded to the N1 (imino) or N2 (amino) nitrogen atom of guanine suggesting that T:G has two metastable states. The first state has two, the latter, less probable state has one hydrogen bond between the two bases. This two-state behaviour is also represented in the two free energy minima of shift and shear. The base pair opening angle exhibits a second free energy minimum, too, which is located at a significantly higher angle than the first minimum or the free energy minima of the Watson-Crick pairs.

The molecular dynamics simulations of the T:G mispair show significant local distortions comparable to those found by NMR experiments [6]. We observe λ -angles similar to those reported in [6] for all three base pairs (cf. Figure S4). However, in our simulations the mispair has a second state with a much higher λ -angle. Moreover, the wobble pair is kinetically unstable and fluctuates between two states, one of which is closer to canonical B-form conformation than the other. The more distorted state is less probable and as such can be regarded as only transiently occupied. This second state has not been observed in the structures of T:G containing DNA, modelled from solution NMR data. This discrepancy can be due to overestimation of the open state by the force field used in the simulations and/or underestimation due to the restraints applied in the structure modelling based on the

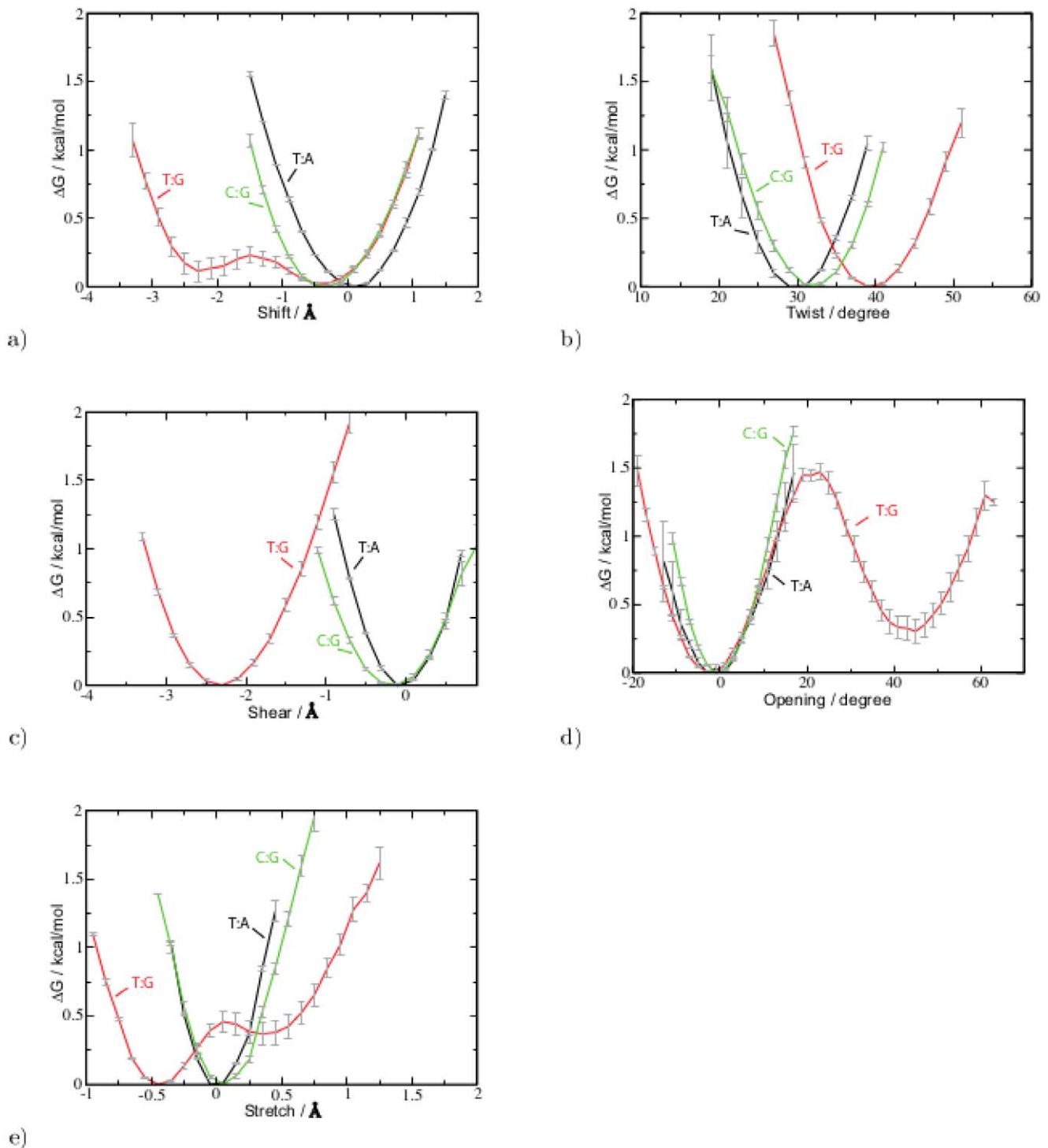


Figure 3. Free energy profiles of local DNA parameters of the T:G mismatch (red) and C:G (green) and T:A (black) pair, respectively, obtained from unbiased MD simulations. Only those parameters with significant differences are shown a) shift b) twist c) shear d) opening e) stretch. For the free energy profiles of the other parameters see Figures S2 and S3 in the supplementary material.
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NMR spectra. The occasional (partial) openings of the base pair as observed in our simulations may be sufficient to be detected by the searching glycosylase and induce further inspection by the enzyme such as transitions from an “interrogating complex” to an “excision complex” [5] including flipping the base.

The free energy profile computed for the base flipping of thymine or cytosine out of T:G, T:A or C:G pairs, respectively, shows that the mismatched thymine is the most probable to reach an extra-helical conformation. The energy required for the single base flip (5 kcal/mol) is about the same as computed previously for the combined rotation of both, G and T, opening through the

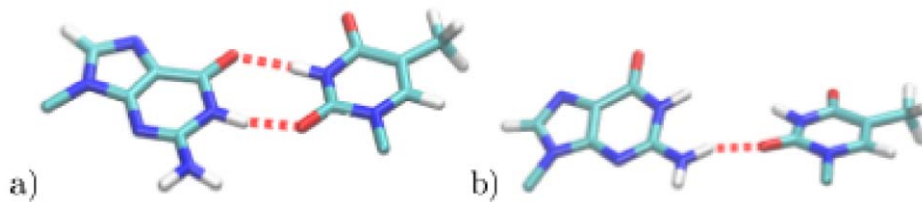


Figure 4. Snapshots of the unbiased simulation of T:G. a) Two-hydrogen bonds conformation, b) one-hydrogen bond conformation.

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major groove [12]. The free energy cost for flipping a single T out of a T:A pair (10 kcal/mol) and C out of a C:G pair (11 kcal/mol) are similar to each other but significantly higher than that computed for mispaired thymine. The error range for the cytosine flip in the higher flip angle regime, however, is significantly larger than for thymine from T:A pairs. This is due to higher energy pathways having been sampled and also indicates a larger conformational heterogeneity of the flipped-out cytosine. Note that the free energy profile shown in Fig. 6 is computed from averaging over several independently calculated free energy profiles. In case of the T:G mispair these individual runs differ from each other in terms of the exact location of the two free energy minima and the barrier between them (cf. Figure S20). This variance indicates that another conformational change, taking place on longer time scales, has not been fully averaged out in the individual runs. However, the similarity between the profiles of the flip angle obtained for the unbiased and the averaged ABF simulations suggests that by averaging over several individual runs the insufficient sampling of the individual runs could be compensated to some extent. Among the many possible “slow degrees of freedom” a flip and re-stacking of the complementary base are the most probable candidates. We have observed such transitions occasionally in some runs, which are not included in the present analysis.

In other computational studies of single base flips from C:G and T:A pairs similar to the one presented here, varying results are reported. Depending on the surrounding sequence, the force field, and opening restraint applied the energetic cost for flipping thymine (from T:A) has been calculated to about 13 kcal, and 15–22 kcal have been computed for flipping cytosine (in C:G) [10,11,28]. Despite the variation of the detailed numbers in all the computational studies, the single base flip of mispaired T

requires less energy than base flip of a pyrimidine from T:A or C:G Watson-Crick pairs. This is in agreement with the order of equilibrium constants for base pair opening, obtained from experimental imino proton exchange rates [65], which are larger by two orders of magnitude in T:G than in T:A and by one order of magnitude in T:A than in C:G.

However, as has been pointed out previously [10–12], imino proton exchange can take place already at an opening angle of approximately 30° from equilibrium (i.e. 70° flip angle). Our results are in agreement with this finding, showing that at a flip angle of 70° both, solvent accessibility of the imino proton (N3-H) and the number of its hydrogen bonds to water, increase significantly. In case of the mispaired T, the conformation with a 70° flip angle is populated even without the application of an external force allowing hydrogen bonds between the imino proton and solvent water to be observed. This would explain the unusually long life time of the “open state” (as determined by proton exchange kinetics) reported in [12]: The partially-open state with 70° flip angle is a second, metastable state of the T:G wobble pair. In the Watson-Crick pairs, 70° flip angle conformations are not stable and are about 6 kcal/mol higher in energy and as such unlikely to be observed. The partially open/flipped-out state (70° flip angle) clearly shows how the dynamics of the DNA is changed due to the G:T mispair.

One can speculate that the intrinsic dynamics of mispaired DNA plays a role in discriminating by the enzyme. The partially-open state, observed in our simulations, would serve as an indicator of mispaired T in G:T, as opposed to A:T which could be recognised by the repair enzyme in a more passive mechanism: The Glycosylases which process T:G mispairs first recognise local distortions in the base steps and base-pair geometries which deviate from normal B-form DNA. Moreover, a partially open

Table 2. Occupancies of hydrogen bonds between DNA base pairs C:G, T:A, and mispair T:G, computed from the simulation of paired and mispaired DNA.

paired (C:G)			mispair (T:G)			paired (T:A)		
Donor	Acceptor	Occ./%	Donor	Acceptor	Occ./%	Donor	Acceptor	Occ./%
G-N2	C-O2	90 ± 2	G-N2	T-O2	34 ± 6	A-N6	T-O4	78 ± 1
G-N1	C-N3	91 ± 4	G-N1	T-O2	72 ± 3			
C-N4	G-O6	68 ± 5	T-N3	G-O6	50 ± 6	T-N3	A-N1	96 ± 1
water	C-O2	68 ± 29	water	T-O2	17 ± 3	water	T-O2	65 ± 3
C-N4	water	56 ± 4	water	T-O4	71 ± 3	water	T-O4	57 ± 9
			T-N3	water	35 ± 7			

Hydrogen bonds of C and T with bulk water are listed, too. G and A are the corresponding bases on the complementary strand in the C:G and T:A pair, respectively. For atom labels see Figure 2.

doi:10.1371/journal.pone.0053305.t002

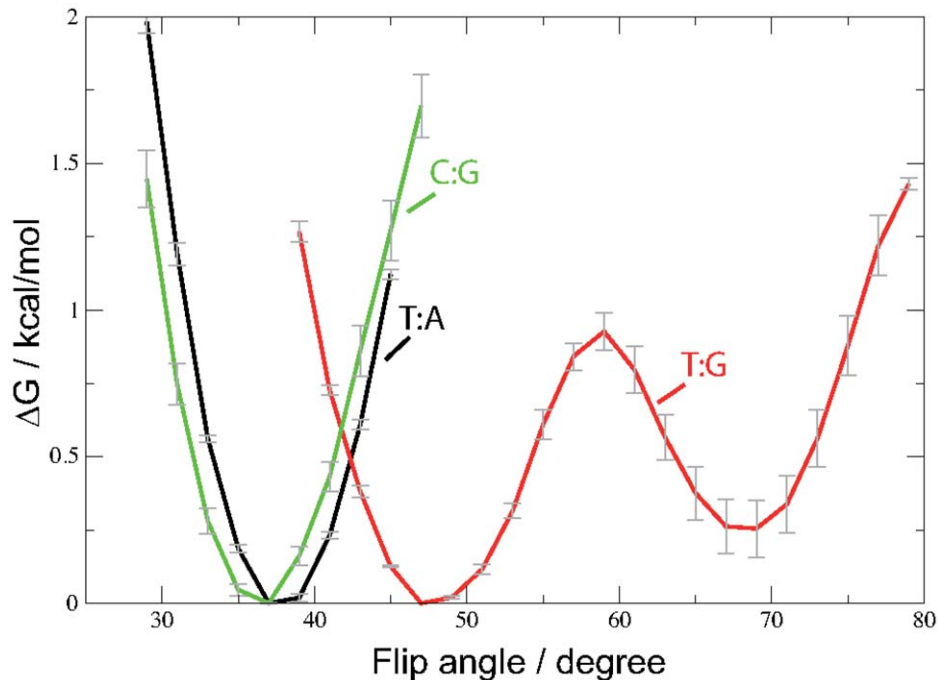


Figure 5. Free energy profiles of the pseudo-dihedral flip angle evaluated from the unbiased MD simulations of the T:G mispair (red) and C:G (green) and T:A (black) pair, respectively.
doi:10.1371/journal.pone.0053305.g005

state of the T:G mispair, which we observe to be transiently occupied also in the unbiased simulations, is supposedly easy to be recognized by the searching repair enzyme.

Recognition of the helical distortions as exhibited by the mispaired T:G, as opposed to T:A, and subsequent formation of a tight “interrogative” [5] protein-T:G complex, may help to save the enzyme from processively trying to flip each base and thereby

also avoid flipping of a paired (T:A) thymine and to erroneously remove it.

Conclusion

DNA containing a single T:G mispair exhibits local dynamics significantly different from DNA without such a mispair. The T:G wobble pair shows a distorted conformation compared to T:A or C:G pairs. Moreover, besides the completely intra-helical state, it

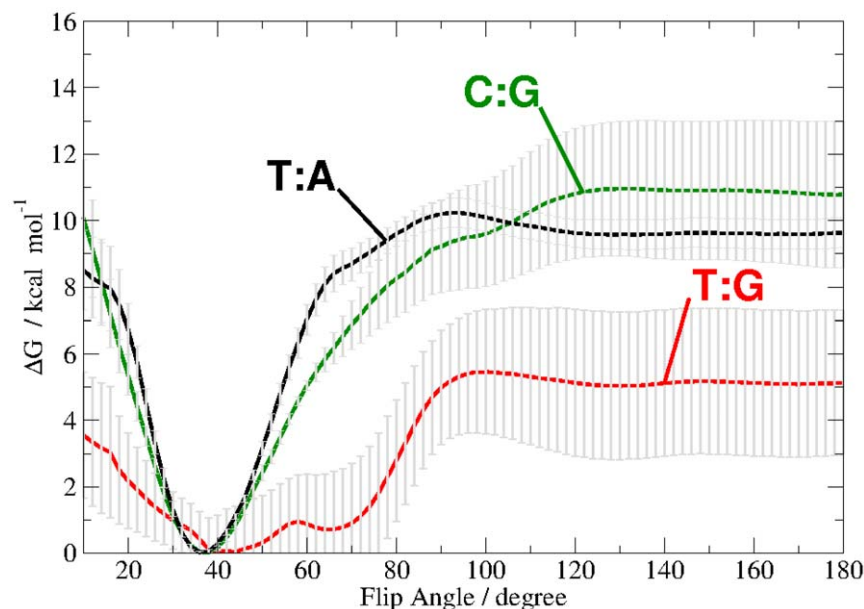


Figure 6. Free Energy profile of the base flip for thymine of a T:G mispair (red), cytosine of a C:G pair (green) or thymine of a T:A pair (black). The pseudo dihedral coordinate is illustrated in Figure 1.
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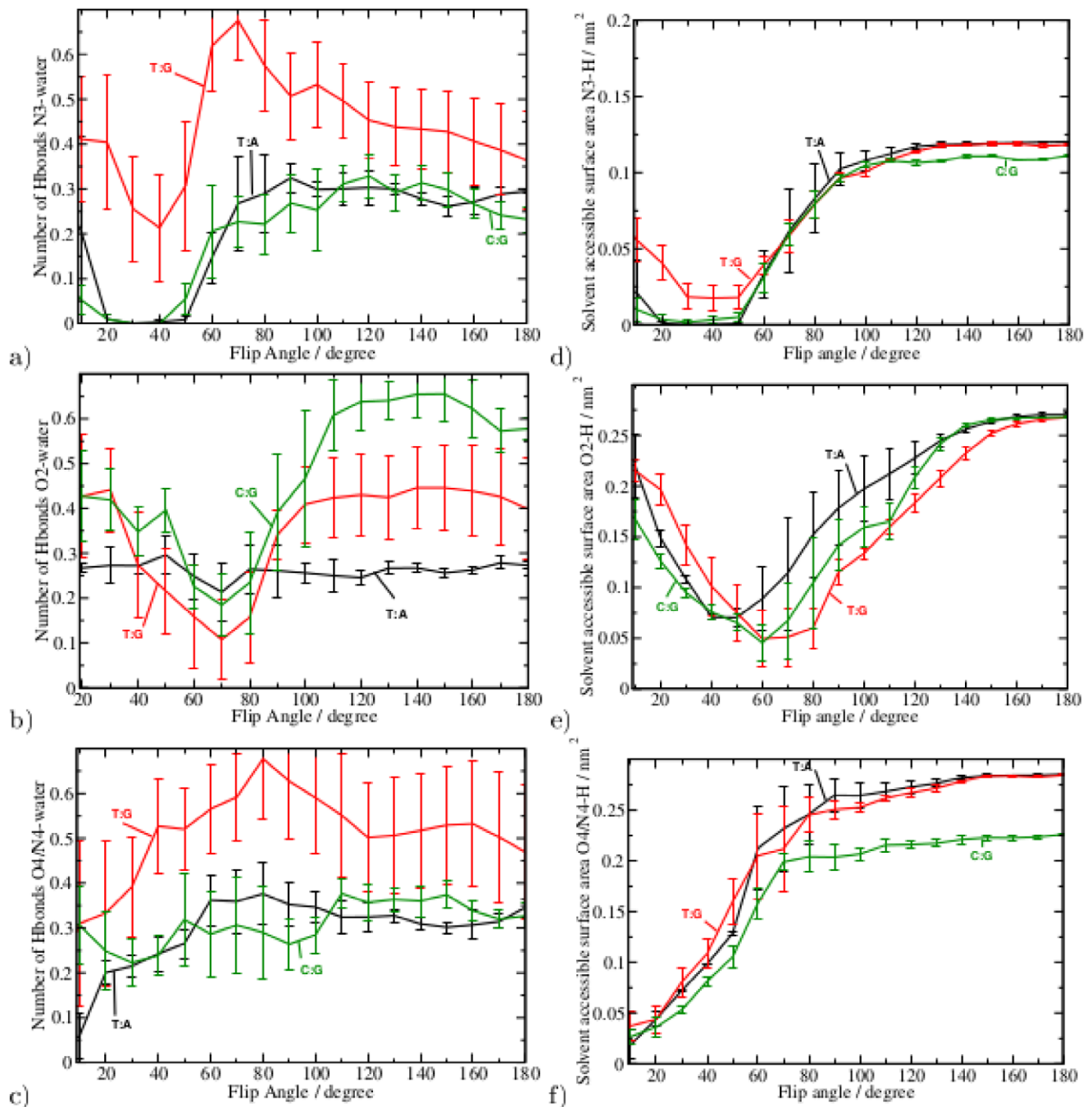


Figure 7. Left: Average number of Hydrogen bonds of solvent water with base atoms a) N3-H b) O2 and c) O4 or N4H2 in case of thymine or cytosine, respectively, as a function of the base flip angle. Right: Solvent accessible surface area of base atoms d) N3-H e) O2 and f) O4 or N4H2 in case of thymine or cytosine, respectively, as a function of the base flip angle. Data from the simulation of the C:G and T:A pairs are plotted in black and green, respectively, data for the T:G mismatch is shown in red. For atom labels see Figure 2. doi:10.1371/journal.pone.0053305.g007

exhibits a second, less probable metastable state which is partially open/flipped-out and allows the thymine imino proton to be accessed by the solvent water.

Our free energy calculations show that thymine is much more probable to be flipped than cytosine in a C:G pair or thymine in a T:A pair, a fact that could possibly be exploited by the repair enzymes.

Supporting Information

Figure S1 RMSD of the DNA with T:G mismatch (red) and C:G (green) and T:A (black) pair, respectively, as a function of the simulation time. (EPS)

Figure S2 Free energy profiles of local base step parameters of the T:G mismatch (red) and C:G (green) and T:A (black) pair,

respectively, as a function of the simulation time: a) roll b) tilt, and c) twist, d) rise, e) shift, f) slide.

(EPS)

Figure S3 Free energy profiles of local base pair parameters of the T:G mispair (red) and C:G (green) and T:A (black) pair, respectively: a) buckle b) opening, c) propeller, d) shear, e) stretch, and f) stagger.

(EPS)

Figure S4 Free energy profiles of the λ -angles computed from the unbiased MD simulations. a) λ -angle at T or C, respectively, b) λ -angle at A or G of the complementary strand.

(EPS)

Figure S5 Free energy profiles of the twist angle of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S6 Free energy profiles of the tilt angle of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S7 Free energy profiles of the roll angle of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S8 Free energy profiles of rise of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S9 Free energy profiles of slide of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S10 Free energy profiles of shift of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) step 12, b,g,m) step 11, c,i,o) step 10, d,j,p) step 9 and e,k,q) step 8.

(EPS)

Figure S11 Free energy profiles of opening of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

(EPS)

Figure S12 Free energy profiles of propeller twist of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

References

- Lindahl T (1982) DNA repair enzymes. Annual review of biochemistry 51: 61–87.
- Sancar A, Sancar GB (1988) DNA repair enzymes. Annual review of biochemistry 57: 29–67.

(EPS)

Figure S13 Free energy profiles of buckle of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

(EPS)

Figure S14 Free energy profiles of stagger of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

(EPS)

Figure S15 Free energy profiles of shear of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

(EPS)

Figure S16 Free energy profiles of stretch of the T:G mispair (red, a–e) and T:A (black, f–k) and C:G (green, l–q) pair, respectively, at different simulation times. a,f,l) base pair 12, b,g,m) base pair 11, c,i,o) base pair 10, d,j,p) base pair 9 and e,k,q) base pair 8.

(EPS)

Figure S17 Free energy profiles of the flip angle of a) the T:G mispair (red), b) T:A (black) and c) C:G (green) pair, respectively, at different simulation times.

(EPS)

Figure S18 Time series of the flip angle in T:G, computed from the unbiased MD simulations.

(EPS)

Figure S19 PMF of the Flip Angle at different simulation times.

(EPS)

Figure S20 Free energy profile of individual ABF simulations of the base flip in T:G.

(EPS)

Figure S21 Snapshots of the DNA base flip simulation of C:G (top), T:G (middle), and T:A (bottom), taken at (left) the flipped-in state (free energy minimum), (middle) at 60–70° flip angle and (right) at the flipped-out state.

(EPS)

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Author Contributions

Conceived and designed the experiments: PI. Performed the experiments: PI. Analyzed the data: PI MZ. Contributed reagents/materials/analysis tools: PI MZ. Wrote the paper: PI MZ.

- Stivers JT, Jiang YL (2003) A mechanistic perspective on the chemistry of DNA repair glycosylases. Chemical reviews 103: 2729–2760.
- Stivers JT (2008) Extrahelical damaged base recognition by DNA glycosylase enzymes. Chemistry-A European Journal 14: 786–793.

5. Friedman JI, Stivers JT (2010) Detection of damaged DNA bases by DNAGlycosylase enzymes. *Biochemistry* 49: 4957–4967.
6. Allawi HT, SantaLucia, Jr J (1998) NMR solution structure of a DNA dodecamer containing single G-T mismatches. *Nucl Acid Res* 26: 4925–4934.
7. Leroy JL, Kochoyan M, Huynh-Dinh T, Gueron M (1988) Characterization of base-pair opening in deoxynucleotide duplexes using catalyzed exchange of the imino proton. *J Mol Biol* 200: 223–238.
8. Gueron M, Leroy JL (1995) Studies of base pair kinetics by nmr measurement of proton exchange. *Methods Enzymology* 261: 383–413.
9. Kochoyan M, Gueron M, Leroy JL (1987) A single mode of dna base-pair opening drives imino proton exchange. *Nature* 328: 89–92.
10. Banavali NK, MacKerell, Jr AD (2002) Free energy and structural pathways of base flipping in a DNA GCGC containing sequence. *J Mol Biol* 319: 141–160.
11. Giudice E, Varnai P, Lavery R (2003) Base pair opening within B-DNA: free energy pathways for GC and AT pairs from umbrella sampling simulations. *Nucl Acid Res* 31: 1434.
12. Varnai P, Canalia M, Leroy JL (2004) Opening Mechanism of G-T/U Pairs in DNA and RNA Duplexes: A Combined Study of Imino Proton Exchange and Molecular Dynamics Simulations. *J Am Chem Soc* 126: 14659–14667.
13. Arora K, Schlick T (2004) In Silico Evidence for DNA Polymerase- β 's Substrate-Induced Conformational Change. *Biophys J* 87: 3088–3099.
14. Beveridge DL, Barreiro G, Byun KS, Case DA, Cheatham, III TE, et al. (2004) Molecular dynamics simulations of the 136 unique tetranucleotide sequences of DNA oligonucleotides. I. Research design and results on d(CpG) steps. *Biophys J* 87: 3799–3813.
15. Chen J, Dupradeau FY, Case DA, Turner CJ, Stubbe JA (2008) DNA oligonucleotides with A, T, G or C opposite an abasic site: structure and dynamics. *Nucleic Acids Research* 36: 253.
16. Dalhus B, Laerdahl JK, Backe PH, Bjorås M (2009) DNA base repair-recognition and initiation of catalysis. *FEMS microbiology reviews* 33: 1044–1078.
17. Dixit SB, Beveridge DL, Case DA, Cheatham, III TE, Giudice E, et al. (2005) Molecular dynamics simulations of the 136 unique tetranucleotide sequences of DNA oligonucleotides. II: sequence context effects on the dynamical structures of the 10 unique dinucleotide steps. *Biophys J* 89: 3721–3740.
18. Fadda E, Pomès R (2011) On the molecular basis of uracil recognition in DNA: comparative study of TA versus UA structure, dynamics and open base pair kinetics. *Nucl Acid Res* 39: 767.
19. Fujii S, Kono H, Takenaka S, Go N, Sarai A (2007) Sequence-dependent DNA deformability studied using molecular dynamics simulations. *Nucl Acid Res* 35: 6063.
20. Hagan MF, Dinner AR, Chandler D, Chakraborty AK (2003) Atomistic understanding of kinetic pathways for single base-pair binding and unbinding in DNA. *Proc Nat Acad Sci USA* 100: 13922.
21. Hu J, Ma A, Dinner AR (2008) A two-step nucleotide-flipping mechanism enables kinetic discrimination of DNA lesions by AGT. *Proc Nat Acad Sci USA* 105: 4615–4620.
22. Huang N, Banavali NK, MacKerell, Jr AD (2003) Protein-facilitated base flipping in DNA by cytosine-5-methyltransferase. *Proc Nat Acad Sci USA* 100: 68.
23. Lavery R, Zakrzewska K, Beveridge D, Bishop TC, Case DA, et al. (2010) A systematic molecular dynamics study of nearest-neighbor effects on base pair and base pair step conformations and fluctuations in B-DNA. *Nucl Acid Res* 38: 299.
24. MacKerell, Jr AD, Nilsson L (2006) Theoretical studies of nucleic acids and nucleic acid-protein complexes using CHARMM. *Computational Studies of RNA and DNA* : 73–94.
25. Pérez A, Luque EJ, Orozco M (2007) Dynamics of b-dna on the microsecond time scale. *J Am Chem Soc* 129: 14739–14745.
26. L-Yang, Beard WA, Wilson SH, Roux B, Broyde S, et al. (2002) Local Deformations Revealed by Dynamics Simulations of DNA Polymerase β . with DNA Mismatches at the Primer Terminus. *J Mol Biol* 321: 459–478.
27. Zahran M, Daidone I, Smith JC, Imhof P (2010) The mechanism of DNA recognition by the restriction enzyme *EcoRV*. *J Mol Biol* 401: 415–432.
28. Varnai P, Lavery R (2002) Base-flipping in DNA: pathways and energetic studies with Molecular Dynamics Simulations. *J Am Chem Soc* 124: 7272–7273.
29. Cheatham TE (2004) Simulation and modeling of nucleic acid structure, dynamics and interactions. *Current Opinion in Structural Biology* 14: 360–367.
30. Isaacs RJ, Spielmann HP (2004) Insight into G-T mismatch recognition using molecular dynamics with time-averaged restraints derived from NMR spectroscopy. *J Am Chem Soc* 126: 583–590.
31. Huang N, MacKerell, Jr AD (2004) Atomistic view of base flipping in DNA. *Philosophical Transactions of the Royal Society of London Series A: Mathematical, Physical and Engineering Sciences* 362: 1439–1460.
32. Zacharias M (2006) Minor groove deformability of DNA: A molecular dynamics free energy simulation study. *Biophys J* 91: 882–891.
33. Priyakumar UD, MacKerell, Jr AD (2006) Base flipping in a GCGC containing DNA dodecamer: a comparative study of the performance of the nucleic acid force fields, CHARMM, AMBER, and BMS. *J Chem Theor Comput* 2: 187–200.
34. Curuksu J, Zakrzewska K, Zacharias M (2008) Magnitude and direction of DNA bending induced by screw-axis orientation: influence of sequence, mismatches and abasic sites. *Nucleic Acids Research* 36: 2268.
35. Orozco M, Noy A, Pérez A (2008) Recent advances in the study of nucleic acid flexibility by molecular dynamics. *Current opinion in structural biology* 18: 185–193.
36. O'Neil L, Wiest O (2008) Sequence dependence in base flipping: experimental and computational studies. *Org Biomol Chem* 6: 485–492.
37. Lavery R, Zakrzewska K, Beveridge D, Bishop TC, Case DA, et al. (2009) A systematic molecular dynamics study of nearest-neighbor effects on base pair and base pair step conformations and fluctuations in B-DNA. *Nucl Acid Res*.
38. Banavali NK, MacKerell, Jr AD (2002) Free energy and structural pathways of base flipping in a DNA GCGC containing sequence. *J Mol Biol* 319: 141–160.
39. Bouvier B, Grubmüller H (2007) A molecular dynamics study of slow base flipping in DNA using conformational flooding. *Biophys J* 93: 770–786.
40. Priyakumar UD, MacKerell, Jr AD (2006) Computational approaches for investigating base flipping in oligonucleotides. *Chemical reviews* 106: 489–505.
41. Daniels DS, Woo TT, Luu KX, Noll DM, Clarke ND, et al. (2004) DNA binding and nucleotide flipping by the human DNA repair protein AGT. *Nat Struct Mol Biol* 11: 714–720.
42. Szczepanowski RH, Carpenter MA, Czapsinska H, Zaremba M, Tamulaitis G, et al. (2008) Central base pair flipping and discrimination by PspGI. *Nucl Acid Res* 36: 6109.
43. Maiti A, Morgan M, Pozharski E, Drohat A (2008) Crystal structure of human thymine DNA glycosylase bound to DNA elucidates sequence-specific mismatch recognition. *Proc Nat Acad Sci USA* 105: 8890–8895.
44. Maiti A, Morgan M, Drohat A (2009) Role of two strictly conserved residues in nucleotide flipping and N-glycosylic bond cleavage by human thymine DNA. *J Biol Chem* 284: 36680–36688.
45. Maiti A, Noon MS, MacKerell Jr AD, Pozharski E, Drohat AC (2012) Lesion processing by a repair enzyme is severely curtailed by residues needed to prevent aberrant activity on undamaged DNA. *Proc Nat Acad Sci USA* 109: 8091–8096.
46. Maiti A, Drohat A (2011) Dependence of substrate binding and catalysis on ph, ionic strength, and temperature for thymine DNA glycosylase: Insights into recognition and processing of gt mispairs. *DNA repair* 10: 545–553.
47. Brooks BR, Bruccoleri RE, Olafson BD, States DJ, Swaminathan S, et al. (1983) CHARMM: A program for macromolecular energy, minimization and dynamics calculations. *J Comput Chem* 4: 187–217.
48. Wang J, Cieplak P, Kollman P (2000) How well does a restrained electrostatic potential (resp) model perform in calculating conformational energies of organic and biological molecules? *J Comput Chem* 21: 1049–1074.
49. Pérez A, Marchán I, Svobiz D, Sponer J, Cheatham III TE, et al. (2007) Re-nement of the AMBER force field for nucleic acids: improving the description of $[\alpha, \gamma]$ conformers. *Biophys J* 92: 3817–3829.
50. MacKerell, Jr AD, Banavali N, Foloppe N (2000) Development and current status of the CHARMM force field for nucleic acids. *Biopolymers* 56: 257–265.
51. Jorgensen WL, Chandrasekhar J, Madura JD, Impey RW, Klein ML (1983) Comparison of simple potential functions for simulating liquid water. *J Chem Phys* 79: 926–935.
52. Darden T, York D, Pedersen LG (1993) Particle mesh Ewald: an Nlog(N) method for Ewald sums in large systems. *J Chem Phys* 98: 10089–10092.
53. Evans DJ, Holian BL (1985) The Nose-Hoover thermostat. *J Chem Phys* 83: 4069.
54. Ryckaert JP, Cicotti G, Berendsen HJC (1977) Numerical integration of the cartesian equations of motion of a system with constraints: molecular dynamics of n-alkanes. *J Comp Phys* 23: 327–341.
55. Darve E, Pohorille A (2001) Calculating free energies using average force. *J Chem Phys* 115: 9169–9183.
56. Chipot C, Héning J (2005) Exploring the free-energy landscape of a short peptide using an average force. *J Chem Phys* 123: 244906.
57. Darve E, Rodríguez-Gómez D, Pohorille A (2008) Adaptive biasing force method for scalar and vector free energy calculations. *J Chem Phys* 128: 144120.
58. Phillips JC, Braun R, Wang W, Gumbart J, Tajkhorshid E, et al. (2005) Scalable molecular dynamics with NAMD. *J Comput Chem* 26: 1781–1802.
59. Eisenhaber F, Lijnzaad P, Argos P, Sander C, Scharf M (1995) The Double Cube Lattice Method: Efficient Approaches to Numerical Integration of Surface Area and Volume and to Dot Surface Contouring of Molecular Assemblies. *J Comput Chem* 16: 273–284.
60. Humphrey W, Dalke A, Schulten K (1996) VMD – Visual Molecular Dynamics. *Journal of Molecular Graphics* 14: 33–38.
61. DeLano WL (2002) The pymol molecular graphics system. DeLano Scientific, San Carlos, CA, USA.
62. Lavery R, Sklenar H (1988) The definition of generalized helicoidal parameters and of axis curvature for irregular nucleic acids. *J Biomol Struct Dyn* 6: 63–91.
63. Ravishanker G, Swaminathan S, Beveridge DL, Lavery R, Sklenar H (1989) Conformational and helicoidal analysis of 30 ps of molecular dynamics on the d(CGCGAATTCGCG) double helix: “curves”, dials and windows. *Journal of biomolecular structure dynamics* 6: 669–699.
64. van der Spoel D, Lindahl E, Hess B, Groenhof G, Mark AE, et al. (2005) GROMACS: fast, flexible, and free. *J Comput Chem* 26: 1701–1718.
65. Moe JG, Russu IM (1992) Kinetics and Energetics of Base-Pair Opening in 5'-d(CGCGAATTCGCG)-3' and a Substituted Dodecamer Containing G:T Mismatches. *Biochemistry* 31: 8421–8428.