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Crystal structures of 2-acetylaminofluorene and 2-aminofluorene in complex with T7 DNA polymerase reveal mechanisms of mutagenesis

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Contributed by Charles C. Richardson, September 22, 2004

The carcinogen 2-acetylaminofluorene forms two major DNA adducts: N-(2′-deoxyguanosin-8-yl)-2-acetylaminofluorene (dG-AAF) and its deacetylated derivative, N-(2′-deoxyguanosin-8-yl)-2-aminofluorene (dG-AF). Although the dG-AAF and dG-AF adducts are distinguished only by the presence or absence of an acetyl group, they have profoundly different effects on DNA replication. dG-AAF poses a strong block to DNA synthesis and primarily induces frameshift mutations in bacteria, resulting in the loss of one or two nucleotides during replication past the lesion. dG-AF is less toxic and more easily bypassed by DNA polymerases, albeit with an increased frequency of misincorporation opposite the lesion, primarily resulting in G → T transversions. We present three crystal structures of bacteriophage T7 DNA polymerase replication complexes, one with dG-AAF in the templating position and two others with dG-AF in the templating position. Our crystallographic data suggest why a dG-AAF adduct blocks replication more strongly than does a dG-AF adduct and provide a possible explanation for the diverse replication properties of these adducts. Moreover, most spectral, enzymatic, and theoretical studies suggest that the AF structure produces much less distortion in the DNA helix than does the closely related AAF adduct (12–16) and that these structural differences are responsible for the different biological consequences.

NMR and molecular mechanics studies of a dG-AAF adduct in double-stranded DNA have confirmed that a major structural distortion is caused by this adduct. These spectral studies have shown that the glycosidic torsion angle of the modified nucleotide adopts a syn conformation with the modified guanine unpaired and lying outside the double helix (17). This distortion of the DNA structure is consistent with an earlier base-displacement model proposed on the basis of circular dichroism studies of the DNA lesion (18) in which the fluorene ring of dG-AAF partially inserts between adjacent base pairs, causing a local disruption of base-pairing. This model is termed the insertion–denaturation model (19, 20).

Studies of the deacetylated dG-AF adduct in double-stranded DNA by similar methods have revealed two interchangeable conformers of the modified nucleoside (21–24). In one conformer, the fluorene ring of dG-AF is oriented outside the DNA helix, where it does not perturb base-pairing. In this outside-binding conformation, the modified guanosine adopts an anti conformation and base-pairs with the complementary DNA strand. In the alternative conformer of dG-AF, the fluorene ring stacks inside the double helix, and the guanosine adopts a syn conformation in which it is exposed and unpaired in a manner similar to the dG-AAF adduct.

The highly distorting syn conformation of the dG-AAF adduct has been used to explain why this modification poses a strong block to replication and is not efficiently copied by a polymerase (11). It is possible that the ability of the dG-AAF adduct to halt replication may also promote frameshift mutagenesis by providing sufficient time for the realignment of primer and template DNA strands to form a slipped mispair with the lesioned nucleotide looped out of the template strand (25). Similarly, the syn conformation of dG-AF would presumably interfere with base-pairing in the active site of a DNA polymerase, whereas the

Abbreviations: AAF, 2-acetylaminofluorene; dAAF, N-(2′-deoxyguanosin-8-yl)-AAF; dG-AAF, N-(2′-deoxyguanosin-8-yl)-2-acetylaminofluorene; dG-AF, N-(2′-deoxyguanosin-8-yl)-2-aminofluorene; ddATP, 2′,3′-dideoxyadenosine triphosphate; ddCTP, 2′,3′-dideoxycytidine triphosphate.

Data deposition: The atomic coordinates and structure factors have been deposited in the Protein Data Bank, www.pdb.org [PDB ID codes IX95 (dG-AAF complex), IX95 (dG-AF complex grown in the presence of ddCTP), and IX95 (dG-AF complex grown in the presence of ddATP)].

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suggesting that the lesion does not support base-pairing interactions within the polymerase active site (10, 32). Furthermore, the substrate-induced conformational change of the polymerase is not observed when nucleotides are added to the complex of the Klenow fragment bound to DNA containing a dG-AAF adduct (30). Additional support for a distorted conformation of dG-AAF-modified DNA in complex with a DNA polymerase comes from footprinting studies of T7 DNA polymerase in complex with native and AAF-lesioned DNAs (33). In contrast to a dG-AAF lesion, DNA containing a dG-AAF adduct binds the Klenow fragment less strongly, and the polymerase can undergo a conformational change upon the addition of dCTP (30).

The different binding modes of DNA templates containing a dG-AAF or dG-AF adduct to a DNA polymerase are likely to account for the strikingly different mutagenic spectra caused by these lesions. Here we present crystal structures of the replicative T7 DNA polymerase (consisting of the gp5 catalytic subunit and its processivity factor, E. coli thioredoxin) in complex with dG-AAF- and dG-AF-modified DNA substrates that reveal the basis for the strong blockage of DNA synthesis by a dG-AAF adduct and suggest why bypass of a dG-AF adduct can occur with a high probability of misinsertion opposite the lesion.

Methods
Preparation of Modified DNAs and Replication Complexes. A 26-mer template strand containing a single dG was designed (5′-CCCGATCACAATACATCTCC-3′), along with a complementary primer strand (5′-GGAGAGTATTGG-TAGTGTGA-3′), and both oligonucleotides were chemically synthesized. N-acetoxy-AAF was obtained from the National Cancer Institute repository operated by the Midwest Research Institute (Midland, TX). The 26-mer template strand and 21-mer primer strand were purified by electrophoresis on denaturing polyacrylamide gels. The template strand was reacted with N-acetoxy-AAF to form a single dG-AAF adduct and then purified by reverse-phase HPLC (34). A portion of the dG-AAF oligonucleotide was deacetylated by incubating in 1 M sodium hydroxide and 0.25 M 2-mercaptoethanol for 2 h at 37°C. The resulting DNA containing dG-AAF was purified again by HPLC.

An exonuclease-deficient T7 DNA polymerase (Δ118–123 T7) was purified as described in ref. 35. Replication complexes were assembled with either the dG-AAF- or dG-AF-containing DNA template by incubating T7 DNA polymerase (0.1 mM protein) with a stoichiometric amount of the DNA template and primer strands in buffer containing 0.5 mM 2′,3′-dideoxythymidine triphosphate (ddTTP) and 10 mM incoming nucleotide, either 2′,3′-dideoxycytidine triphosphate (ddCTP) or 2′,3′-dideoxyadenosine triphosphate (ddATP). Complexes with either ddCTP or ddATP as the incoming nucleotide were separately assembled and crystallized for the dG-AF complex. For the dG-AAF DNA, a binary complex with T7 DNA polymerase lacking an incoming nucleotide also was assembled for crystallization.

Crystallization and X-Ray Structure Determination. Complexes of T7 DNA polymerase with DNA were crystallized under conditions described in ref. 35, and the crystals were harvested in the presence or absence of added ddCTP or ddATP and then frozen. X-ray diffraction data were collected at station F1 of the Cornell High Energy Synchrotron Source (CHESS) (Ithaca, NY). The x-ray data were indexed, integrated, and scaled by using the HKL2000 package (36), and the structures were determined by the molecular replacement method by using the program EPMR (37). A model of a native ternary complex of T7 DNA polymerase (Protein Data Bank ID code 1T7P) was positioned in the unit cell after nucleotides within the polymerase active site and residues from the surrounding fingers subdomain were removed. These regions around the active site were rebuilt into the omit electron density calculated from the partial model of the complex, and the resulting crystallographic models were refined by...
Table 1. X-ray data collection and refinement statistics

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<th>Complex</th>
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<th>dG-AF/ddATP</th>
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*Values in parentheses are for the last resolution shell.
†R = ∑(|Fo| – |Fc|)/|Fo|, where |Fo| and |Fc| are observed and calculated structure factor amplitudes. Rmerge is calculated with 5% of randomly selected reflections that were not used in refinement.

Using the program CNS (38) (Table 1). The coordinates for the dG-AAF complex (PDB ID code 1X9M), the dG-AF complex grown in the presence of ddCTP (PDB ID code 1X9S), and the dG-AF complex grown in ddATP (PDB ID code 1X9W) have been deposited in the Protein Data Bank.

Results

Structure of the dG-AAF-Containing Replication Complex. When dG-AAF is in the templating position of a replication complex with T7 DNA polymerase, the modified nucleoside is flipped out of the polymerase active site and bound to the surface of the fingers subdomain (Fig. 2). The syn conformation of the dG-AAF nucleoside (Fig. 2c) facilitates the insertion of the fluorene ring into a hydrophobic pocket behind the O-helix of the fingers subdomain (Fig. 2d). The fluorene adduct stacks against Phe-528 and other aliphatic side chains that are normally buried in the core of the fingers subdomain (Fig. 2 b and d and Fig. 4, which is published as supporting information on the PNAS web site). The acetyl group of dG-AAF is not contacted by the polymerase and lies outside the hydrophobic pocket with the oxygen atom facing solvent. The N2 amino group of the guanine base donates a hydrogen bond to Asp-534, and N7 of guanine accepts a hydrogen bond to Arg-566 (Fig. 2d). These electrostatic contacts with guanine further stabilize the interaction of the DNA adduct with the fingers.

In comparison with a native ternary replication complex (35), T7 DNA polymerase in complex with dG-AAF-modified DNA adopts an open, distorted conformation of the fingers. This change in conformation involves a hinge-like rotation of the fingers subdomain away from the polymerase active site and repacking of helices within the fingers in response to the intercalation of the dG-AAF nucleoside. The fingers are rotated ~35° away from the fully closed position that is observed in complexes with native DNA and a bound nucleotide (39). There is no evidence for a bound ddCTP molecule in the crystal structure of the dG-AAF complex, despite our having grown the crystals in the presence of 10 mM ddCTP. In fact, repacking of the fingers subdomain caused by the insertion of the AAF adduct

Fig. 2. dG-AAF adopts the syn conformation with the fluorene ring intercalated into the polymerase fingers domain. (a) dG-AAF lies outside the polymerase active site, and the fingers domain of the polymerase is in an open conformation. The T7 gene 5 protein is shown as ribbons, with the O-helix within the fingers highlighted in red. The primer strand of the DNA is in light red, the template strand in yellow, and ddG-AF is in cyan and green. The region around the dG-AAF binding site is circled. (b) Enlarged view of the circled region in a. The fluorene ring (green) of dG-AAF (cyan and green) is inserted into the polymerase fingers domain. (c) The simulated annealing omit electron density around the region of the syn dG-AAF of the dG-AAF-containing complex is shown in stereo, contoured at 2.5 σ above the mean value. (d) Interaction between dG-AAF and the protein. Hydrophobic side chains (gold) of the fingers form a pocket around the fluorene ring. Two charged residues (pink), Arg-566 and Asp-534, form hydrogen bonding interactions with the G base.
be superimposed well on structures of T7 DNA polymerase in complex with an unmodified DNA and a nucleotide.

**Structure of a dG-AF-Containing Replication Complex.** The overall structure of T7 DNA polymerase in complex with DNA containing a dG-AF adduct is very similar to that of the dG-AAF complex described above (Table 1 and Fig. 3), but there are two notable differences. First, electron density corresponding to the dG-AF nucleotide is absent from several different structures determined by using crystals grown in the presence of ddCTP (2.7-Å resolution) or ddATP (2.4-Å resolution) (Table 1; see Methods for crystallization conditions). Although the structure of the protein and the bound DNA is well defined by the x-ray experiment, the electron density around the dG-AF adduct is of limited quality. However, the crystal structure clearly shows that the dG-AF nucleoside does not bind in the polymerase active site and that the side chain of Tyr-530 occupies the binding site for the template base in a closed ternary complex. Sketchy electron density on the surface of the polymerase in the dG-AF complex is suggestive of the 5′ end of the template strand extending away from the normal binding site for the template base. However, the fluorene ring of the dG-AF adduct is not intercalated into the hydrophobic pocket on the surface of the fingers where the syn dG-AAF nucleotide binds. In the dG-AF structure, the surrounding nonpolar side chains rotate inward to fill the pocket.

The fingers of the polymerase in the dG-AF complex are in an open conformation, closely resembling their overall position in complexes with dG-AAF DNA, and, likewise, there is no evidence of a bound nucleotide for crystals grown in the presence of 10 mM ddCTP or ddATP (Fig. 3). Harvesting crystals in the continued presence of ddCTP (5 mM) also had no effect on the structure. It is surprising, therefore, that crystals of the dG-AF complex grew only in the presence of ddCTP or ddATP. This observation suggests that the incoming nucleotide stabilizes the polymerase–DNA complex during some stage of the crystallization experiment but that the open conformation of the polymerase lacking bound nucleotide is energetically preferred.

We have modeled the anti conformation of the dG-AF nucleoside in the active site of the closed conformation of T7 DNA polymerase by superimposing the modified guanine on the crystal structure of a native dG–ddCTP complex (40). By rotating about the C8–N2 bond, the fluorene ring of the bound dG-AF adduct can be positioned so that it does not clash with the closed fingers of the polymerase. However, after the nucleotide binds to the templating position, rotation about the C8–N2 bond would be restricted by clashes between the N2 proton and the sugar phosphate backbone of the dG-AF nucleoside, as well as by interference between the fluorene ring and neighboring protein groups as the nucleotide transits through unfavorable conformations. In the syn conformation, the aromatic fluorene ring of the dG-AF adduct might favorably stack with the base pair adjacent to the polymerase active site, effectively competing with binding of the guanine template in the anti conformation seen in native complexes (35, 40). These restrictions on positioning the nucleotide adduct for base-pairing interactions with the incoming nucleotide may explain the observed pause in DNA synthesis opposite dG-AF (29).

**Discussion**

**dG-AAF and dG-AF Interfere with Nucleotide Binding and Induced Fit.** T7 DNA polymerase and other polymerases undergo a conformational change during nucleotide incorporation in which the fingers subdomain opens and closes by a hinge-like motion (31). The open conformation allows the active site to sample nucleotides until the fingers close around a nucleotide correctly paired with a template base (35, 41). Bulky lesions such as a cis-syn cyclobutane thymine dimer in a DNA template shift the con-
formational equilibrium toward the open conformation of the fingers, slowing DNA synthesis (42). The fingers of T7 DNA polymerase are in an open conformation in complexes with dG-AAF and dG-AF templating DNA synthesis, and there is an additional distortion of the O-helix (Fig. 3 and Fig. 5, which is published as supporting information on the PNAS web site).

NMR studies of double-stranded DNA containing a dG-AAF adduct show that it exists predominantly in a syn conformation with its fluorene ring intercalated between base pairs (17). In complex with T7 DNA polymerase, the dG-AAF nucleoside maintains the syn conformation, but the adduct binds in a surprising location, inserting in a hydrophobic pocket behind the O-helix of the fingers subdomain. This mode of interaction with the polymerase halts DNA synthesis by locking the fingers in an open conformation (Fig. 2) and causing the Tyr-530 side chain to encroach on the nucleotide binding site (Fig. 3). Interca
telation of the fluorenyl moiety of dG-AAF causes a repacking of hydrophobic residues within the core of the fingers subdomain (Fig. 2c), distorting the C terminus of the O-helix. This conformational change brings Tyr-530 into the active site of the polymerase and prevents nucleotide substrates from binding (Fig. 3). This shift of Tyr-530 might explain the strong block of dG-AF incorporation across from the modified nucleoside. Moreover, Tyr-530 described here is more direct. In the native ternary complex of T7 DNA polymerase, Tyr-530 stacks against the Tyr-714 residue (Tyr-714) of Bacillus stearothermophilus DNA polymerase in complex with a thymine photodimer, the lesioned base is swung out of the helix (42). Here, this residue fills in the "pseudoabasic site" created when the bulky thymine dimer adduct cannot be accommodated in the polymerase active site. The role of Tyr-530 described here is more direct. In the native ternary complex of T7 DNA polymerase, Tyr-530 stacks against the templating base that pairs with the incoming nucleotide (35).

Structural Insights into the Mutagenic Potentials of dG-AAF and dG-AF. In bacteria, there is a preference for a dG-AAF adduct to cause frameshift mutations and for a dG-AF adduct to cause base substitution mutations, mostly G → T transversions (4). The different mutagenic outcomes of replication bypass of dG-AAF and dG-AF lesions in DNA have been ascribed to differences in the internal conformation and flexibility of these modified nucleosides (Fig. 1), with dG-AAF causing greater distortion of DNA structure than dG-AF (18, 25, 26, 45). Our structural studies of these adducts in the context of a DNA polymerase reported here verify some aspects of this model and offer additional insights about a surprising mechanism for inducing replication blockage.

Single-turnover kinetic studies show that T7 DNA polymerase inserts dCMP opposite dG-AF and dG-AAF ~104- and 106-fold more slowly, respectively, than it does opposite an unmodified dG (29). dAMP is misincorporated across from dG-AAF adducts, albeit at a slower rate than that at which dCMP is incorporated, whereas dAMP is the preferred nucleotide opposite a dG-AAF (29). In the syn conformation of dG-AF, the guanine base would lie outside the templating position, where it is unable to pair with an incoming nucleotide. This situation would resemble synthesis past an abasic site, which causes DNA synthesis to pause and favors the eventual insertion of dAMP (46, 47). A similar situation could also hold for incorporation opposite a dG-AAF adduct in the syn conformation, allowing it to slowly misincorporate dAMP across from the modified nucleoside. However, insertion by the A-rule would likely require the fluorenyl moiety of dG-AAF to dissociate from the polymerase so that the fingers could close and assist with positioning, activating the (unpaired) nucleotide for incorporation. In contrast to dG-AF, a dG-AAF nucleoside adopts the anti conformation infrequently, and it supports templated DNA synthesis at a vanishingly small rate. Thus, it may appear that there is a preference for dAMP incorporation across from dG-AAF.

Residues participating in the hydrophobic interactions that stabilize dG-AF binding in the fingers subdomain are located far away from the polymerase active site, yet they play a major role in blocking the enzyme function. However, the role of Tyr-530 described here is more direct. In the native ternary complex of T7 DNA polymerase, Tyr-530 blocks against the templating base that pairs with the incoming nucleotide (35). The analogous residue (Tyr-671) of the Thermus aquaticus DNA polymerase plays the same role, acting as a chaperone or "positioning device" for the templating base in the polymerase active site (41). In crystal structures of T7 DNA polymerase complexed to a thymine photodimer, the lesioned base is swung out of the helix (42). Here, this residue fills in the "pseudoabasic site" created when the bulky thymine dimer adduct cannot be accommodated in the polymerase active site. The analogous residue (Tyr-714) of Bacillus stearothermophilus DNA polymerase is reported to act as a flexible chaperone that guides the transport of primer-template base pairs through the polymerase during DNA synthesis (48). The chaperoning function of Tyr-530 takes an interesting turn in the dG-AAF- and dG-AF-containing complexes reported here, where the O-helix is pushed toward the active site by the adduct and the side chain of Tyr-530 blocks the binding of nucleotides. Removal of this bulky side chain could relieve the blockage and perhaps facilitate untemplated bypass of dG-AAF. In fact, replacement of the analogous residue in the Klenow fragment polymerase (Tyr-766) with a smaller serine residue outside the polymerase active site might affect the conformation of the fingers subdomain, leading to aberrant polymerase behavior.
The binding of dG-AAF on the surface of the fingers subdomain provides several key elements for the formation of a slipped mispair that is compatible with deletion frameshift mutagenesis during lesion bypass (25). First, the stable binding of the DNA template outside the polymerase active site dramatically slows DNA synthesis, providing time for the primer strand to realign with the lesioned nucleotide looped out of the template. Second, the fingers are held in an open conformation that imposes fewer restraints on the 3′ end of the primer, allowing it to dissociate and establish alternative base-pairing interactions with the template. Finally, the interactions of the polymerase with the dG-AAF nucleotide interfere with or prevent base-pairing to the lesion while keeping the DNA anchored to the polymerase. These factors increase the likelihood of forming a slipped mispair that excludes the lesioned nucleotide from the duplex while preventing dissociation of the DNA substrate from the polymerase during lesion bypass.

In summary, it has long been suspected that the structural difference between the dG-AAF and dG-AF adducts in duplex DNA somehow were correlated to the differences in replicative and mutagenic properties of these adducts in cellular systems. However, it was not clear how the structures that formed in double-stranded DNA are related to the structures at the primer-template junction bound in the active site of DNA polymerase before and during nucleotide insertion opposite the lesion. The crystal structures reported here provide a molecular basis for the observed effects of these DNA adducts on replication and reveal a correspondence between the structures of the adducts in the polymerase active site and those in duplex DNA.

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