The N-clasp of human DNA polymerase $\kappa$ promotes blockage or error-free bypass of adenine- or guanine-benzo[a]pyrenyl lesions

Lei Jia$^1$, Nicholas E. Geacintov$^2$ and Suse Broyde$^{1,2,*}$

$^1$Department of Biology and $^2$Department of Chemistry, New York University, 100 Washington Square East, New York, NY 10003, USA

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ABSTRACT

DNA bypass polymerases are utilized to transit bulky DNA lesions during replication, but the process frequently causes mutations. The structural origins of mutagenic versus high fidelity replication in lesion bypass is therefore of fundamental interest. As model systems, we investigated the molecular basis of the experimentally observed essentially faithful bypass of the guanine 10S-(+)-trans-anti-benzo[a]pyrene-$N^2$-dG adduct by the Y-family human DNA polymerase $\kappa$, and the observed blockage of pol $\kappa$ produced by the adenine 10S-(+)-trans-anti-benzo[a]pyrene-$N^2$-dA adduct. These lesions are derived from the most tumorigenic metabolite of the ubiquitous cancer-causing pollutant, benzo[a]pyrene. We compare our results for the dG adduct with our earlier studies for the pol $\kappa$ archaeal homolog Dpo4, which processes the same lesion in an error-prone manner. Molecular modeling, molecular mechanics calculations and molecular dynamics simulations were utilized. Our results show that the pol $\kappa$ N-clasp is a key structural feature that accounts for the dA adduct blockage and the near-error-free bypass of the dG lesion. Absence of the N-clasp in Dpo4 explains the error-prone processing of the same lesion by this enzyme. Thus, our studies elucidate structure-function relationships in the fidelity of lesion bypass.

INTRODUCTION

Bulky lesions normally block the progress of processive high fidelity DNA polymerases (1–5). Subsequently, the replicative polymerase may be replaced by one or more translesion bypass polymerases (6–10). The error-free bypass of DNA lesions during translesion synthesis (TLS) is a repair pathway, which plays an important role in coping with highly blocking lesions in all three domains of life (6,10). However, translesion synthesis is often carried out by low-fidelity bypass DNA polymerases (11–14). In error-prone TLS, the mismatched nucleotides incorporated opposite the lesion introduce mutations into the genome that can result in the development of cancer (15,16). In humans, there are four known Y-family bypass DNA polymerases, pol $\eta$, pol $\iota$, pol $\kappa$ and Rev1 (7,17). Pol $\kappa$ is the only Y-family polymerase found in humans which has homologs in both bacteria (DinB or Pol IV in Escherichia coli) and archaea (Dbh in Sulfolobus acidocaldarius and Dpo4 in Sulfolobus solfataricus) (18).

The structural features of low-fidelity bypass polymerases that allow for TLS of some lesions but not others, while high-fidelity polymerases are stalled by the same lesions, is a topic of significant interest (3,9,10). The structures of a number of Y-family polymerases have been investigated (10,19). A ternary complex crystal structure of pol $\kappa$ lacking its C-terminal domain has recently been determined (20). This structure contains the major domains of pol $\kappa$ including its signature N-terminal domain, namely, the ‘N-clasp’, an unmodified DNA primer/template complex, and an incoming dTTP Watson–Crick paired with the templating base adenine (20). The N-clasp in pol $\kappa$ is at present unique, as it has not yet been observed in any other Y-family polymerase. With the N-clasp on the major groove side (Figure 1) of the nascent duplex, pol $\kappa$ completely encircles a region of the DNA containing the template–primer junction at its active site. It is noteworthy that pol $\kappa$ lacking the N-clasp (pol $\kappa_{68-526}$) is severely impaired in its DNA synthesis activity (20).

The ability of pol $\kappa$ to synthesize DNA both in the absence and presence of lesions has received considerable attention. In the case of unmodified DNA templates, pol $\kappa$ synthesizes DNA with low fidelity (21), and it has also been shown that it is promiscuous in extending primer strands past mispaired primer–template termini (22). The processing of DNA lesions by pol $\kappa$, whether error-free or error-prone, has been shown to be highly lesion...
Benzo[a]pyrene (B[a]P) is a ubiquitous cancer-causing environmental pollutant, present in tobacco smoke and as a product of fossil fuel combustion (16). The most mutagenic and tumorigenic metabolite of B[a]P is the (+)-7,8-dihydrodiol-9,10-epoxy-7,8,9,10-tetrahydrobenzo[a]pyrene, (+)-anti-B[a]PDE (39,40). The (+)-anti-B[a]PDE molecule is unstable in aqueous environments and reacts readily with the exocyclic amino groups of adenine or guanine in DNA by the predominant trans opening of the epoxide, to form the 10S(+)-trans-anti-B[a]P-N6-dA and –N2-dG adducts (B[a]P-dA and B[a]P-dG, respectively) (41) (Figure 2A). These adducts are the major adenine and guanine adducts (41–43) and have been extensively investigated in primer extension studies with a number of replicative and bypass polymerases (2,5,33–36), or by a C-terminal truncated form of pol κ (37), with mostly correct insertion of dCTP opposite these lesions, and is thus error-free. Pol κ has also been implicated in the error-free bypass of these benzo[a]pyrene-derived DNA adducts in mouse embryonic cells (51), and in mouse embryo fibroblasts transfected with a gapped plasmid containing a site-specifically incorporated benzo[a]pyrene-derived N2-dG adduct (52).

Primer extension studies by Rechkoblit et al. (33) using the (+)-trans-B[a]P-N2-dG and –N6-dA adducts, employing pol κ as the polymerase, have shown particularly interesting differences, namely, error-free bypass of B[a]P-dG and essentially full blockage of the polymerase by B[a]P-dA adducts under similar experimental conditions. Specifically, pol κ incorporates dCTP opposite the B[a]P-dG adduct, although at a slowed rate relative to unmodified template strands. However, the progress of pol κ is strongly impeded by the B[a]P-dA adduct, and it stalls at the step that involves nucleotide insertion opposite the lesion (33). The structural and mechanistic factors underlying this striking difference in primer extension efficiency are of great interest because they provide us with further insights into the abilities of Y-family polymerases to bypass some DNA lesions but not others. The availability of the crystal structure of pol κ (20) provides an excellent opportunity for examining the structural features of this polymerase that allow for the TLS of B[a]P-dG adducts, but not of the B[a]P-dA adducts, utilizing molecular modeling and dynamics approaches.

Our hypothesis is that the unique pol κ N-clasp plays an important role in the differential treatment of these bulky N2-dG and N6-dA adducts. To investigate this hypothesis, we modeled these B[a]P-dG and B[a]P-dA adducts on the templating strands in the pol κ active site region in two positions shown in Figure 1. The first position is termed the preinsertion position: in this case an unmodified template A. The second position we studied is the insertion position, in which the dNTP is paired with the B[a]P-dG or the B[a]P-dA adduct. For this position, we modeled dCTP paired with
the dG adduct or dTTP paired with the dA adduct (Figure 2B). We then carried out an extensive automated van der Waals (VDW) energy survey with 373, 248 different conformations for each lesion; this produced a comprehensive topography of the energy as a function of the lesion conformation within the polκ ternary complex. The survey was employed to locate structures with minimum steric contacts and thereby to provide initial structures for subsequent molecular dynamic (MD) simulations. Our results reveal that the N-clasp plays a critical role in the blockage by the B[a]P-dA adduct, which competes for space with the N-clasp and is without steric close contacts. Moreover, at the insertion position, the B[a]P ring system is not near the N-clasp and is always in close contact with it when in the preinsertion position. However, in the case of the dG adduct in this position, the B[a]P ring is near the active site, and favors slippage in the bypass of this adduct.

Figure 2. (A) Structures of the 10S(+)trans-anti-B[a]P-N6-dA and the 10S(+)trans-anti-B[a]P-N6-dG adducts. The absolute configurations of the four chiral atoms C7, C8, C9 and C10 are indicated. Torsion angles are defined as $\chi = O4'-(dR)-C1'$ (dR = deoxyribose), $\chi = N1-C6-N^\delta-C10(B[a]P-dA)$ or $N1-C2-N^\delta-C10(B[a]P-dG)$, $\chi = C6-N^\delta-C10(B[a]P-dA)-C9(B[a]P-dA)$ or $C2-N^\delta-C10(B[a]P-dG)-C9(B[a]P-dG)$. (B) Base sequences of the four models. The incoming nucleotide dNTP is indicated in blue font. In the case of the dA adduct in the insertion position, dTTP was employed in the VDW energy survey model, while dATP was employed in the MD simulations (see text).

METHODS

Construction of the initial models by molecular modeling

We utilized the ternary crystal structure of polκ with DNA and incoming dTTP (PDB ID: 2OH2, complex B) (20) as the template to build our initial models. This polκ structure did not resolve 57 residues (residue 225–281) in its palm subdomain. These residues are not present in the polκ homolog Dpo4 and are called the flexible ‘dorsal tendril’ (55). We superimposed this polκ complex with a Dpo4 ternary complex (PDB ID 2AGQ) (56) and found that the two enzymes share high structural similarity in the palm domain regardless of the unresolved residues in the polκ structure (Supplementary Figure S1). Moreover, these residues are not located near the active site, and modeling them could lead to a problematic structure. Therefore, to obtain a reasonable model of a continuous polκ structure, we closed the gap between residues 224 and 282 as a loop and discarded the missing 57 residues; the program MODELLER running at the ModLoop server was employed to model the loop containing residues 223, 224, 282 and 283 (57,58). Furthermore, this polκ structure contains only one Mg2+ in its active site; however, a functional polymerase requires two Mg2+ ions to organize the active site (59,60). The nucleotide binding Mg2+ ion (MgA), already present in the structure, has normal octahedral coordination (61,62). We modeled the
missing catalytic Mg\(^{2+}\) (MgB) (Supplementary Figure S2) in the pol \(\kappa\) active site following octahedral coordination criteria from a high resolution pol \(\beta\) crystal structure and a computational study with Dpo4 (61,62). The backbone torsions of the primer nucleotide were modestly adjusted (<30\(^\circ\)) in order to achieve the coordination distance between O3\(^\prime\) and the catalytic Mg\(^{2+}\). Coordinates of all initial structures are provided in Supplementary data.

The DNA sequences were then modified to the ones used by Rechkoblit et al. (33) in conducting the primer extension studies. These sequences are shown in Figure 2B. To complete our initial models, the B[\(\omega\)]P-dA and -dG adducts were modeled at the preinsertion and insertion positions (Figure 1) by covalently linking the lesion, taken from an NMR solution structure of this adduct in an 11-mer B-DNA duplex (63), to the amino groups of adenine or guanine. Templating bases were Watson–Crick partnered with their complementary dNTPs by replacing the original thymine of dTTP in the crystal structure with the appropriate base (Figure 2B), without changing the glycosidic torsion. Four initial models (B[\(\omega\)]P-dA and B[\(\omega\)]P-dG adducts in preinsertion and insertion positions) were created for the energy survey.

Van de Waals energy surveys of the dA and dG adducts in pol \(\kappa\)

To obtain a VDW energy surface of pol \(\kappa\) containing the dA and dG adducts, we carried out an energy survey for the four pol \(\kappa\) models, utilizing the two torsion angles at the linkage site between the damaged base and B[\(\omega\)]P, \(\alpha\) and \(\beta\), and the glycosidic linkage \(\chi\) (Figure 2A), as variables. A torsion driver program was employed to rotate these dihedral angles at 5\(^\circ\) intervals in combination over the whole 360\(^\circ\). Thus, we created \(72^3 = 373,248\) structures for each model. The energy calculations were carried out by the SANDER module in AMBER 9.0 (64) with the Cornell et al. (65) force field and PARM99 (66), ff99SB (67) and parmbsc0 (68) parameter sets. The partial charges for the B[\(\omega\)]P-dG and -dA adducts were parametrized using quantum mechanical ESP calculations (HF/6-31G\(^*\), Gaussian 03) and RESP for fitting the charge to each atom center (69,70). The partial charges for the dNTPs and Mg\(^{2+}\) are taken from Perlow et al. (71,72). All of the added force field parameters, atom types, and topology assignments for the lesions, dNTPs and Mg\(^{2+}\) are listed in Supplementary Data (Tables S1–S7). In these single-point VDW energy calculations, we utilized an implicit solvent model with dielectric constant represented by the function \(\varepsilon = 4r\) (\(r\) is the distance between an atom pair). TECPLOT10 from Amtec Engineering Inc., was employed to generate 3D energy maps.

Molecular dynamics simulations of pol \(\kappa\) complexes

From our 3D VDW energy survey maps, we selected the lowest energy pol \(\kappa\) structure of each model as the initial structure for the MD simulations. The same force field was employed as in the energy survey study. In addition, we simulated a normal G • C pair control model without B[\(\omega\)]P modification utilizing the crystal structure (PDB ID: 2OH2) with sequence remodeled to that of the B[\(\omega\)]P-dG adduct in the insertion position (Figure 2B). Detailed MD simulation protocols are given in Supplementary Material. The number of counterions and water molecules added to the system and the sizes of the solvation boxes are given in Supplementary Table S8.

Stability of the molecular dynamics simulations

Plots of the time-dependent root-mean-square deviations (RMSD) of each ensemble member’s binding pocket relative to the first structure of production MD are given in Supplementary Figure S3; the binding pocket is defined as any atom within 5 Å of the damaged nucleotide. This binding pocket for the dA adduct in the preinsertion position is unstable due to local crowding. The binding pocket generally fluctuates in a stable manner after 3.0 ns for the remaining three models. Our analyses were confined to the 5.0–8.0 ns time frame as a uniform window for all simulations.

Structural analyses

Snapshots of the structures during the simulations with solvent and counterions removed were obtained with the PTRAJ module of the AMBER 9.0 suite (64). We selected one best representative structure from each ensemble for graphic illustration. To pick the best representative structure, we extracted 100 structures at 30 ps intervals from the 5.0–8.0 ns time frame for each model. Each extracted structure was superimposed to the remaining 99 structures and the RMSDs were computed and summed. The structure with the smallest sum RMSD was selected as the most representative structure (Supplementary Table S9). This procedure essentially follows the philosophy in MoiView (73), which, however, could not be utilized directly due to the size of our system. PTRAJ was also employed to determine the time-dependence of the RMSD, and the torsion angles \(\alpha\), \(\beta\) and \(\chi\) (Supplementary Figure S4). Detailed hydrogen bonding analyses were carried out with the CARNAL module in AMBER7.0. Criteria for hydrogen bonding are: donor heavy atom to acceptor heavy atom distance \(\leq 3.4\) Å; donor heavy atom-hydrogen-acceptor heavy atom angle: 135\(^\circ\) to 225\(^\circ\). To analyze the reaction-ready active site organization, we measured the Mg\(^{2+}\) octahedral coordination distances (Supplementary Figure S2), the in-line attack distance (O3\(^\prime\)-Pz), and angle (O3\(^\prime\)-Pz-O3z). To compute the occupancy, we established criteria, based on near reaction-ready pol \(\beta\) and Dpo4 structures (61,62), for a well-organized active site: in-line attack distance <3.5 Å; in-line attack angle 180 ± 30\(^\circ\); Mg\(^{2+}\) coordination distance <2.4 Å, except for the distance MgA-O1z: for this distance we accepted <3.0 Å, based on our simulation for the unmodified control and because one long distance has also been observed in several of our earlier simulations (47,74,75). The near reaction ready occupancy is the percent of time during the 5–8ns time frame of the simulation utilized for analysis that meets all these criteria simultaneously (Donny-Clark,K. Shapiro,R. and Broyle,S., submitted).

INSIGHT 2005 from Accelrys, Inc. and PyMOL 1.1 from DeLano Scientific LLC. were utilized for molecular modeling and visualization, respectively. Clustalw2 (76) running at Uniport servers, http://beta.uniprot.org.
RESULTS

The objective of our work was to structurally rationalize the origins of the observed blockage of the dA adduct, while the dG adduct is bypassed with near-high fidelity. We carried out a comprehensive molecular mechanics VDW energy survey for each adduct within the preinsertion and insertion positions of the polymerase–DNA complexes in order to search for feasible domains for the α', β' and χ flexible torsions (Figure 2A) that govern the lesion orientation (see Methods section). Structures with minimal steric collisions then provided initial models for subsequent MD simulations. Full energy spectra, in two-dimensional slices, of the results presented below are given in Supplementary Movies S1–S12.

The molecular mechanics energy surveys reveal that the dA adduct in the preinsertion position has close contacts with the pol κ N-clasp, while the dG adduct does not.

We determined 3D VDW energy topographies for the lesion at the pol κ preinsertion position (Figure 1). The VDW energy was computed for each conformer to evaluate steric hindrance between the lesion and the protein/DNA complex. Three-dimensional (α', β' and χ) energy contour maps for each model were then constructed based on the energy survey. Figure 3 shows the contour maps for these models. Energies are relative to the lowest energy conformer of each model (Supplementary Table S11). Our purpose here was to locate domains that were not severely forbidden. Therefore, at this stage, we visualized structural domains to an energy cutoff of 1000 kcal/mol to locate all potentially feasible structures, including those that could become viable with small protein movements; the steric term in the Lennard–Jones potential for the VDW energy increases very steeply, small protein movements; the steric term in the Lennard–Jones potential for the VDW energy increases very steeply, small torsional movements can produce very large VDW energy increases. For the preinsertion position models, the maps reveal that the dA adduct is high energy essentially throughout the entire torsional space. There is only one domain which is not forbidden by our criteria. This domain, which contains the lowest energy structure, is extremely small (Figure 3, inset). However, the energy map for the dG adduct reveals two much larger such domains (Figure 3 and Supplementary Table S12).

Figures 4A and B show the lowest energy structures of the dA and dG adducts in the pol κ preinsertion position. In the case of the dA adduct, the lowest energy conformation places the adduct in the anti conformation of the glycosidic torsion. However, there are still close contacts between the B[a]P ring system and the pol κ N-clasp (Figure 4), as the B[a]P rings are directed toward it. In the case of the dG adduct, the lowest energy conformation also has the adduct in the anti-conformation. Furthermore, while the B[α]P rings are linked to the N2 of dG, which is on the minor groove side of the guanine in duplex B-DNA, here the adduct is in a single-stranded region; this single-stranded region has a non-B-DNA backbone conformation, which places the B[α]P moiety on the major groove side of the evolving duplex (Figure 4B). There are no close contacts between the adduct and the enzyme, since the B[α]P rings are not near to the N-clasp. In addition, the aromatic ring of Phe29 on the N-clasp stacks with the modified guanine base better than it does with the modified adenine; this stacking helps stabilize the dG adduct in the preinsertion position (Figure 4).

The molecular mechanics energy surveys show that the dG adduct in the insertion position favors the Watson–Crick capable anti-glycosidic conformation but the dA adduct prefers the syn domain.

Here, we were motivated to elucidate the experimentally observed error-free insertion of dCTP opposite the dG adduct as well as the absence of dTTP incorporation opposite the dA adduct (33). The 3D energy maps (Figure 3) show relatively low energy domains for both dA and dG adducts in the pol κ insertion position, but the dG adduct prefers the anti-glycosidic bond conformation while the dA adduct favors the syn domain.

In the case of the dG adduct, the lowest energy anti-conformation is preferred by at least 89 kcal/mol over any syn conformers. The lowest energy structure places the dG adduct in the normal anti-conformation opposite the Watson–Crick partner dCTP, and positions the B[α]P in the pol κ on the minor groove side of the evolving DNA duplex, not near to the N-clasp. In addition, the B[α]P ring system stacks with Phe151 (Figure 4D). The B[α]P ring orientation for the dG adduct is different from that observed in NMR solution structures of this adduct in a full duplex (64), and a model template–primer complex containing a C opposite the lesion (77). In these structures, the B[α]P rings are oriented 5' along the modified strand in the minor groove. In the present pol κ structures, the B[α]P rings are 3' directed along the modified strand. While this is not energetically favored in solution (78), it is enforced in pol κ because steric collisions with the enzyme finger domain would occur with a 5' orientation.

For the dA adduct, we also modeled the normal Watson–Crick partner, dTTP, as the incoming nucleotide. However, the energy survey revealed that the anti-glycosidic bond conformation is prohibited, and hence Watson–Crick pairing is impossible. On a structural level, this stems from the N9 linkage of the adenine to the B[α]P moiety which places the B[α]P ring system on the major groove side of the adducted adenine (Figure 4C); on this side the B[α]P rings collide with the enzyme active site or N-clasp or adjacent DNA and dNTP, depending on the α' and β' linkage site conformations. However, lower energy domains are obtained for the syn glycosidic region (Figure 3).

MD simulations of the lesions in the preinsertion position show crowding by the N-clasp in the dA but not the dG adduct.

The energy survey provided lowest energy conformations as the initial structures for subsequent MD simulations. The MD simulations of the models with lesions in the
preinsertion position show that the dG adduct can be housed in this site stably, as was indicated by the VDW energy survey. The dG adduct is not positioned near to the N-clasp, but is accommodated in the space between the N-clasp and the DNA (Figure 5B). In addition, the stacking interaction between the damaged guanine and Phe29 observed in the energy survey remains in place to help stabilize the adduct (Figure 5B). For the dA adduct, the MD simulation shows slight movement of the adduct and the N-clasp just sufficient to alleviate the close contacts (Figure 5A) between them that were observed in the energy survey. The B[a]P ring system continues to be directed toward the N-clasp during the simulation and is housed in the crowded environment created by it (Figure 6). There is some stacking between Phe29 and the modified adenine, but the stacking interactions are less favorable than with the dG adduct (Figure 5).

The RMSD plots (Supplementary Figure S3) of the active site region containing all atoms that are within 5 Å of any atom of the added nucleotide show that the dA lesion is unstable in the preinsertion position, while the dG adduct is stable. We also analyzed the time dependence for the adduct linkage site torsions \( \alpha' \) and \( \beta' \), and the glycosidic torsion \( \chi \) in the ensemble of structures. The ensemble average values and the SD for these torsion angles are listed in Supplementary Table S12, and the time dependences are given in Supplementary Figure S4. All torsions, but most notably the glycosidic torsion \( \chi \) of
the dA adduct have larger SD values than those of the dG adduct, showing that glycosidic torsional motions contribute significantly to the instability of the dA adduct.

The experimental data revealed that Watson–Crick paired nucleotide incorporation occurred at the unmodified bases 3' to the dG and dA adduct in the preinsertion position (Figure 1). We therefore evaluated the near reaction-ready active site organization for the incoming dNTPs paired with these unmodified bases. Accordingly, we analyzed the Mg2+ coordination, the in-line attack distance (O3'-Pz) and angle (O3'-Pz-O3z) from the MD ensemble derived from the ternary complex, utilizing criteria for ideal active site organization involving Mg2+ coordination, in-line attack distance and attack angle which must be simultaneously met, as detailed in the Methods section. We obtained a near reaction-ready occupancy (the percent of time that the criteria are met) of 83.3% and 62.7% for the dG and dA adduct cases, respectively. By comparison, an unmodified control simulation had an occupancy of 71.1%. In addition, the dNTP is well aligned in a Watson–Crick paired orientation with the unmodified partner cytosine or adenine (Figure 5C and D, Supplementary Table S13). These results suggest that dNTP insertion is feasible when the adduct is in the preinsertion position for both the dG and dA adducts; this is considered further in relation to the experimental data in the Discussion section.

**Figure 5.** MD simulation preinsertion position models for the dA and dG adducts. The best representative structures from the MD ensembles are shown. The color scheme is the same as in Figure 4. Views of (A) the dA adduct and (B) the dG adduct looking into the major groove side of the nascent DNA duplex. The adducts are positioned on the major groove side in both cases. Note the oval which highlights the crowding between the adduct and the N-clasp. Phe29 stacks partially with the damaged adenine but stacks very well with the damaged guanine. The (C) and (D) orientations are obtained from (A) and (B) through a ~90° rotation to obtain views looking approximately down the DNA helix axis; these views show clearly the Watson–Crick base pairing alignment between the purple unmodified base 3' to the lesion and its partner dNTP. Stereoviews are shown in Supplementary Figure S9. Movies are shown in Supplementary data.

**MD simulations of the lesions in the insertion position show that the dG adduct is well accommodated with Watson–Crick pairing to dCTP, while dNTP incorporation is disfavored in the dA adduct**

To further investigate the essentially error-free dCTP incorporation observed opposite the dG adduct, we modeled the adduct at the insertion position partnered with dCTP. The dG adduct forms a Watson–Crick base pair with the incoming dCTP (Figure 7D). The B[a]P ring system is stably accommodated in a shape complementary pocket on the evolving minor groove side of the DNA duplex (Figure 8). The stacking interaction between the B[a]P and Phe151, observed in the energy survey, remains stable during the simulation (Figure 7D). The active site RMSD and the SD values of the torsions σ', β' and γ also show that the dG adduct is very stable in the insertion position (Supplementary Figures S3 and S4, Table S12). In addition, we obtained a near reaction-ready occupancy for the dG adduct of 66.3%.

We wished to determine whether the aromatic Phe151 in pol κ, which stacks with the B[a]P to help stabilize its position for Watson–Crick pairing, is also present in the analogous position of Dpo4. For a broader perspective, we carried out multi-sequence alignment between all four...
human Y-family polymerases, as well as the archaeal enzymes Dpo4 and Dbh with Clustalw2 (76). The results show that only pol κ and Rev1 have a phenylalanine, while other polymerases have a nonaromatic residue in the analogous sequence position: pol η and pol ι have a leucine, and Dpo4 and Dbh have a lysine (Figure 9). These sequence aligned residues are located in analogous structural positions for pol κ and Dpo4, (see Discussion).

To investigate the very small amount of dATP incorporated opposite the dA adduct in the experiments (33), we modeled dATP as the incoming nucleotide for the MD simulation of this adduct (Supplementary Figure S5). The energy survey had revealed that the templating dA adduct favors the syn glycosidic conformation, and this is utilized as the initial model for the MD simulation. Since syn adenine prefers to hydrogen bond with purines in the anti-conformation (79,80), the incoming dATP was placed in the anti-glycosidic conformation (Supplementary Figure S5). Early in the MD simulation, the B[a]P ring system relocated to position the adduct on the DNA major groove side, between the DNA and the N-clasp, and away from the dATP (Figure 7A). This motion, seemingly independent of the dNTP, stems from rotations in the DNA backbone and glycosidic bond torsions (the latter from syn to high anti). This motion appears to be mandated by the proximity of the B[a]P ring system to the 3′-end primer thymine nucleobase in the lowest energy molecular mechanics structure utilized to initiate the MD (Supplementary Figure S6). This movement positions the damaged adenine away from the incoming dATP, thus preventing its hydrogen bonding interactions with any incoming dNTP (Figure 7C, Supplementary Table S13); stacking interactions between the B[a]P-modified adenine and the downstream templating base are also interrupted. The active site RMSD and the SD values of the torsions α', β' and χ, especially χ, show that the dA adduct is less stable than the dG adduct in the

Figure 6. Accommodation of the B[a]P ring system of the dA adduct in the preinsertion position. The adduct in the preinsertion position is directed toward and is crowded by the N-clasp. The color scheme is the same as in Figure 4. The protein is in cartoon mode and the DNA and incoming dNTP are in CPK. (A) The N-clasp is shown in cartoon in order to reveal the position of the B[a]P ring system. The (B) view is the same as (A), except that the N-clasp is shown in CPK. Stereoviews are shown in Supplementary Figure S10.

Figure 7. MD simulation insertion position models for the dA and dG adducts. The best representative structures from the MD ensembles are shown. The color scheme is the same as in Figure 4. Views of (A) the dA adduct and (B) the dG adduct looking into the major groove side of the nascent DNA duplex. The dA adduct is positioned on the major groove side, while the dG adduct is on the minor groove side. The damaged adenine is flipped out of the nascent DNA duplex. The (C and D) orientations are obtained from (A) and (B) through an ~90° rotation to obtain views looking approximately down the DNA helix axis; these views show the alignment between the damaged base and the dNTP. The flipped-out damaged adenine mismatched with dATP cannot hydrogen bond with any dNTP. The damaged guanine is Watson–Crick paired with dCTP and the B[a]P has a stacking interaction with Phe151. Stereoviews are shown in Supplementary Figure S11. Movies are shown in Supplementary data.

Figure 8. CPK model of the B[a]P-dG adduct in the insertion position, looking into the minor groove of the nascent DNA duplex. The color scheme is the same as in Figure 4. The dG adduct is accommodated in a shape-complementary pocket on the evolving DNA minor groove side. The Phe151 residue in magenta stacks with the B[a]P ring system. This MD simulation frame (67591) is among the optimal stacking structures of the ensemble. Stereoviews are shown in Supplementary Figure S12.
insertion position (Supplementary Figures S3 and S4, Table S12). The small amount of dATP incorporation observed in the experiments is considered in the Discussion.

**DISCUSSION**

The pol κ N-clasp plays a central role in determining lesion blockage or bypass

Running start primer extension studies using site specifically modified DNA containing the (+)-trans-B[a]P-N²-dG and N⁰-dA adducts showed that pol κ incorporates mostly dCTP opposite the dG adduct and continues to extend past the lesion. However, for the dA adduct, the polymerase is mainly stopped just before the lesion site; a minor amount of adenine incorporation is observed opposite the adduct after long (30-min) reaction time and there is no further extension. In both the dG and dA adduct cases, nucleotide incorporation opposite unmodified bases 3’ to the lesion is efficient. However, the polymerase stalled after this dNTP incorporation in the dG adduct case. Our modeling studies provide structural insights into these observations.

Well-organized active site for dNTP incorporation opposite the 3’ lesion-flanking base. Our MD simulation of the preinsertion position models (Figure 1B) revealed that the active site is well organized for incorporation of the incoming dNTP opposite the unmodified base 3’ to the lesion, in both the dA and dG adduct cases (Figure 5C and D): Mg²⁺ coordination, O3’ in-line attack distances and angles, and Watson–Crick alignment between the dNTP and its partner base (Supplementary Table S13) are appropriate for a near-reaction ready structure based on criteria derived from a high resolution polymerase crystal structure (61) and QM/MM calculations (62). These results are in agreement with the experimental observation that these dNTP insertions are efficient.

The B[a]P-dA adduct is blocked by the N-clasp at the preinsertion position. Following nucleotide insertion, further progression of the polymerase is mostly blocked by the dA adduct (33,50). The underlying reasons for this blockage are from our energy survey and subsequent MD simulation for the dA adduct in the preinsertion position: our models show that the B[a]P rings are not close to the N-clasp. A non-B-DNA conformation in this single-stranded preinsertion position places the B[a]P rings on the major groove side of the evolving duplex (Figure 5B), while in a full duplex of B-DNA the B[a]P moiety is on the minor groove side (64). Our subsequent studies for the insertion position show that the B[a]P-dG adduct is Watson–Crick-paired with dCTP, with the B[a]P ring system positioned on the minor groove side of the nascent duplex (Figure 7B). There is no steric hindrance in either position. The polymerase pause following nucleotide incorporation opposite the unmodified base 3’ to the lesion in the preinsertion position, may result from the following features: an energy barrier must be overcome to achieve the rearrangement of the lesion from the major to the minor groove side, as it translocates from the preinsertion, to the insertion position. However, once this barrier is overcome, the adduct is well positioned for Watson–Crick pairing of the modified dG with dCTP and for nucleotide incorporation (Figure 7D, Supplementary Table S13). The arrangements of the interacting partners at this active site are well poised for the nucleotidyl transfer reaction, which is consistent with the experimentally observed essentially high fidelity bypass of this lesion.
It is noteworthy that there are other bulky $N^2$-dG adducts that are bypassed by pol $\kappa$, primarily in an error-free manner (32,83,84); these may assume conformations similar to those of the $B[a]P$-$N^2$-dG adduct in the preinsertion and insertion sites of pol $\kappa$. Such adducts include the minor groove conformation of the $N^2$-(2-hydroxyestron-6(\(\alpha,\beta\))-yl)-2'-deoxyguanosine in duplex DNA suggested by a modeling study (85), as well as the 3-(deoxyguanosin-$N^2$)-yl)-2-acetylaminofluorene adduct in double-stranded DNA positioned in the minor groove as shown by NMR solution (86) and modeling studies (87).

**Error-free bypass of the $B[a]P$-$N^2$-dG adduct is supported by specific structural features of pol $\kappa$, notably the N-clasp, not present in the error-prone Dpo4 polymerase**

It is noteworthy that pol $\kappa$ bypasses the $B[a]P$-dG lesion in an essentially error-free manner, while the bypass of the same lesion catalyzed by its DinB prokaryotic homolog Dpo4 is error-prone. In the latter, various mismatches and a prevalence of deletions have been observed, with the sequence context in which the lesion is embedded playing an important role in the specific mutagenic outcome (47,53,54,88). Our modeling and MD simulations for the pol $\kappa$ insertion position provide some structural insights for this striking difference.

In pol $\kappa$, the *anti* glycosidic bond conformation of the damaged templating dG, needed for Watson–Crick pairing at the insertion position, is strongly preferred over the *syn* conformation because the pol $\kappa$ N-clasp has close contacts with the *syn*-oriented $B[a]P$-dG moiety (Figure 10A and B). In contrast, in Dpo4 both *syn* and *anti* $B[a]P$-dG conformations are feasible (47,53,88, Xu, P., Oum, L., Geacintov, N.E. and Broyde, S., manuscript in preparation). The *syn* $B[a]P$-dG adduct is comfortably accommodated on the major groove side, which does not have an N-clasp and is therefore open to the solution environment (Figure 10C). Furthermore, modeling studies show that the *syn* conformation, which is incapable of forming Watson–Crick hydrogen-bonding with the dNTP, can readily support mismatches (47,53,54, Xu, P., Oum, L., Geacintov, N.E. and Broyde, S., manuscript in preparation), thus providing a rationale for the observed low fidelity of Dpo4.

In addition, the aromatic Phe151 residue of pol $\kappa$, positioned in the nascent minor groove of the DNA duplex, stacks with the $B[a]P$ ring system (Figure 8) which helps to stabilize the adducted guanine in an orientation which supports Watson–Crick pairing with dCTP. In contrast, Dpo4 has a lysine residue in the analogous position (Figure 11), which does not contribute to a stabilization of the aromatic ring system. Thus, even when the *anti*-conformation that is capable of Watson–Crick pairing is assumed, in Dpo4 the adducted guanine would be conformationally more flexible and hence more likely to pair with other dNTPs.

**Modeling pol $\kappa$ with bulky DNA lesions: computational considerations**

Finally, we wish to discuss our computational approaches. We utilized molecular dynamics simulations with AMBER 9.0 (64) in order to obtain ensembles of structures to aid in interpreting the experimental data. Simulations of such large systems still pose challenges and require necessary approximations. The first challenge is to obtain initial structures for the subsequent MD
Align function which overlays the backbone Cα atoms. Stereoviews are shown in Supplementary Figure S14.

Figure 11. Superimposition of the anti B[α]P-dG adduct in the pol κ insertion position with the Dpo4 ternary complex (PDB ID: 2AGQ). The most representative structure from the MD ensemble is shown. Pol κ is in gray cartoon; and Dpo4 is in blue cartoon; B[α]P rings are red; the damaged guanine is orange. The magenta Phe151 stacks with the B[α]P ring system. However, the analogous position in Dpo4 is occupied by a green lysine. Superimpositions are performed with the PyMol Align function which overlays the backbone Cα atoms. Stereoviews are shown in Supplementary Figure S14.

simulations. Here, we had to first complete the pol κ crystal structure, which had 57 residues far from the active site and DNA-binding region missing. To avoid possibly serious problems with modeling these residues de novo, we elected to close the gap with a loop as a better option. We also modeled the missing catalytic Mg²⁺ in the active site selected the structure with the lowest VDW energy for each case to initiate the MD simulations with explicit aqueous solvent and Na⁺ counterions. A question arises here: if neighboring structures in the lowest energy region were utilized to initiate the MD, would such structures produce different outcomes in the MD simulations? While the sampling problem in MD simulations (90–92) remains demanding, our sterically very confined pol κ system has extremely limited opportunities for housing the bulky lesions we investigated, and these were ferreted out in the energy survey. Examination of the energy spectra (Figure 3 and Movies S1–S12 in Supplementary Material) reveals the very narrow conformational domains that could be feasible. For the dA adduct, it is very clear that the preinsertion position blocks accommodation of the lesion by collision with the N-clasp, which in and of itself explains the experimental data that this adduct mainly blocks the polymerase. For the dG adduct, the dominant experimental observation to be explained is the virtually error-free incorporation of dCTP opposite the lesion by pol κ. This structural requirement for Watson–Crick base pairing in the enzymatic reaction (see Experimental section) reveals the very narrow conformational domains that could be feasible. For the dA adduct, it is very clear that the preinsertion position blocks accommodation of the lesion by collision with the N-clasp, which in and of itself explains the experimental data that this adduct main...
SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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