Mechanisms of Insertion of dCTP and dTTP Opposite the DNA Lesion \( O^6 \)-Methyl-2′-deoxyguanosine by Human DNA Polymerase \( \eta \)*

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\( O^6 \)-Methyl-2′-deoxyguanosine (\( O^6 \)-MeG) is a ubiquitous DNA lesion, formed not only by xenobiotic carcinogens but also by the endogenous methylyating agent S-adenosylmethionine. It can introduce mutations during DNA replication, with different DNA polymerases displaying different ratios of correct or incorrect incorporation opposite this nucleoside. Of the “translesion” Y-family human DNA polymerases (hpols), hpol \( \eta \) is most efficient in incorporating equal numbers of correct and incorrect C and T bases. However, the mechanistic basis for this specific yet indiscriminate activity is not known. To explore this question, we report biochemical and structural analysis of the catalytic core of hpol \( \eta \). Activity assays showed the truncated form displayed similar misincorporation properties as the full-length enzyme, incorporating C and T equally and extending from both. X-ray crystal structures of both dC and dT paired with \( O^6 \)-MeG were solved in both insertion and extension modes. The structures revealed a Watson-Crick-like pairing between \( O^6 \)-MeG and 2′-deoxythymidine-5′-[\( \alpha,\beta \)-imido]triphosphate (approximating dT) at both the insertion and extension stages with formation of two H-bonds. Conversely, both the structures with \( O^6 \)-MeG opposite dCTP and dC displayed sheared conformation of base pairs but to different degrees, with formation of two bifurcated H-bonds and two single H-bonds in the structures trapped in the insertion and extension states, respectively. The structural data are consistent with the observed tendency of hpol \( \eta \) to insert both dC and dT opposite the \( O^6 \)-MeG lesion with similar efficiencies. Comparison of the hpol \( \eta \) active site configurations with either \( O^6 \)-MeG:dC or \( O^6 \)-MeG:dT bound compared with the corresponding situations in structures of complexes of Sulfolobus solfataricus Dpo4, a bypass pol that favors C relative to T by a factor of \( \approx 4 \), helps rationalize the more error-prone synthesis opposite the lesion by hpol \( \eta \).

High fidelity replication of DNA is essential to all forms of life. However, DNA is extensively modified by both sources outside and inside mammalian bodies. One type of damage is alkylating agents, and the simplest and most common of these are methylyating agents. Methylation of DNA by xenobiotic chemicals, including some used to treat cancer (1), was recognized at least as early as 1961 and then associated with tumors caused by simple alkyl N-nitrosamines (2). Although \( N^7 \)-MeG is the major DNA adduct formed by methylyating agents (2), several lines of evidence indicate that \( O^6 \)-MeG is the most miscoding and mutagenic adduct (3). \( O^6 \)-MeG is formed from endogenous sources as well as exogenous (4), and all mammalian DNA samples show finite levels, generally attributed to extraneous methylation by S-adenosylmethionine (4).

Exactly how \( O^6 \)-MeG miscodes has been a topic of interest ever since it became possible to study miscoding. The general phenomenon with \( O^6 \)-MeG is the insertion of C, the correct base, or T (5). The purines (A, G) have not been incorporated to any major extent by any DNA polymerases studied to date.

Melting studies with oligonucleotides containing \( O^6 \)-MeG provide some (unexplained) results (6). Substitution of \( O^6 \)-MeG for dG lowered the \( Tm \) of a duplex of 12-mers by 19–26 °C. However, the most stable complex was with \( O^6 \)-MeG:C pairing, and the \( O^6 \)-MeG:T pair (favored in misincorporation studies) was the weakest (7, 8). Swann (6) proposed that polymerases either (a) mistake \( O^6 \)-MeG for A due to “physical similarities” and insert dTTP and/or (b) miscoding is a consequence of the alignment of the bases. Both of these explanations are vague, but Swann (6) proposed a wobble pair for \( O^6 \)-MeG:C and Watson-Crick pairing (both with only one H-bond) for \( O^6 \)-MeG:T pairing. These later proved to be the structures observed in the X-ray crystal structure with Sulfolobus solfataricus Dpo4 (with the \( O^6 \)-MeG:T pair being in a two H-bond pseudo-Watson-Crick pairing) (9).

NMR studies of oligonucleotides (in the absence of polymerases) provided evidence for the existence of a wobble base pair

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* The abbreviations used are: MeG, methylguanine; hpol, human DNA polymerase; dCMP-NPP, 2′-deoxycytidine-5′-[\( \alpha,\beta \)-imido]triphosphate; dTMP-NPP, 2′-deoxythymidine-5′-[\( \alpha,\beta \)-imido]triphosphate.

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between O6-MeG and C (10). Crystallographic studies of such oligonucleotides (O6-MeG:C) (no polymerase) yielded split-hydrogen bond bifurcated pairing and Watson-Crick type pairing in Z-DNA (11, 12). NMR studies suggested that O6-MeG:T pairing retained Watson-Crick geometry, with a single hydrogen bond between the N2 exocyclic amine of O6-MeG and the O2 carbonyl of T (10), plus no bonding or possibly a “long” bond between the N1 atom of O6-MeG and the imino proton at the N3 atom of T. A crystal structure of oligonucleotides containing an O6-MeG:T pair (in a self-complementary system) showed Watson-Crick pairing (13).

Three sets of crystal structures of O6-MeG paired with C and T in DNA polymerases are available, and they differ dramatically. With DNA polymerase I of the thermophile Bacillus stearothermophilus, C was paired opposite O6-MeG in an isos-teric Watson-Crick geometry, invoking a rare C tautomer (14). An O6-MeG:dTTP complex in the same enzyme involved an unusual electrostatic interaction between (O6-MeG) O-methyl protons and a dT carbonyl oxygen. S. solfataricus Dpo4 complexes show pairing predicted by NMR studies, i.e. wobble pairing between O6-MeG and C and pseudo-Watson-Crick pairing for O6-MeG and T (9). The only other reported DNA polymerase structure with O6-MeG (and the only one to date with a eukaryotic polymerase) is that of hpol δ, which shows Hoogsteen pairing (generally typical of this enzyme) for both the O6-MeG:C and O6-MeG:T complexes, with one H-bond in the T complex and two H-bonds in the C complex (15). Dpo4 favors C incorporation >T by a factor of ~4 (9), but hpol δ and B. stearothermophilus pol I favor T > C incorporation by factors of ~6 and 10, respectively (14–16). Thus, there appear to be multiple mechanisms by which DNA polymerases catalyze dCTP and dTTP incorporation opposite O6-MeG.

Of the human Y-Family DNA polymerases, hpol η (Uniprot entry Q9Y253, POLH) is by far the most efficient in catalyzing both dCTP and dTTP incorporation (16). This enzyme, like hpol δ, incorporates dCTP and dTTP with similar efficiencies (16). We solved insertion and extension structures with either dCTP (dC) or dTMP-NPP (dT) positioned opposite O6-MeG at the active site of hpol η. The structural data allow insight into the principles underlying the similar activities with both pyrimidine nucleotides displayed by this human TLS (translesion synthesis) polymerase.

Results

Analysis of Misincorporation—Studies with the catalytic core of hpol η and a 23-mer/18-mer template-primer DNA duplex construct with incorporated O6-MeG (Table 1) yielded results essentially identical to our findings with the full-length enzyme (16). The ratio of \( \frac{(k_{cat}/K_m)_{dCTP}}{(k_{cat}/K_m)_{dTTP}} \) was 1.1 (results not presented), identical to the value found earlier (16).

LC-MS analysis of the extended primer indicated the presence of only C and T incorporation opposite O6-MeG (47% C, 53% T) (Fig. 1, Table 2). The ratio is very similar to the results of assays done before with full-length hpol η (16) and in accord with the results of the steady-state kinetic analysis. Thus, C and T are equally preferred.

Crystal Structures of Insertion-stage Ternary hpol η Complexes—We determined two structures of complexes between hpol η and 12-mer template-8-mer primer duplexes (see Table 1 for sequences) containing O6-MeG, with nucleoside triphosphates paired opposite the adducted residue. The atomic coordinates and structure factors have been deposited in the Protein Data Bank under codes 5L1I (insertion O6-MeG opposite dCTP), 5L1J (insertion O6-MeG opposite dTMP-NPP), 5L1K (extension after O6-MeG opposite dC), and 5L1L (extension after O6-MeG opposite dT). In the first, determined to a resolution of 2.79 Å, the incoming nucleotide is dCTP, and in the second, determined to 1.94 Å resolution, the incoming nucleotide is dTMP-NPP. A summary of crystal data, data collection, and refinement parameters is provided in Table 3, and examples of the quality of the final electron density in the active site region are shown in Fig. 2, A and B. Crystals of the complex with the non-hydrolyzable dTTP analog dTMP-NPP were obtained readily, and electron density maps generated after molecular replacement and initial rigid body refinement revealed a fully occupied nucleoside triphosphate molecule. Conversely, dCMP-NPP in combination with Mg2+ in the crystal of the other insertion-stage complex was only partially ordered, with electron density visible for the triphosphate moiety and part of the sugar but not for the base. Subsequently, we resorted to dCTP at higher concentrations in the presence of Ca2+ (to prevent nucleotide insertion) to obtain crystals of the ternary complex in which the entire incoming nucleotide was surrounded by electron density (Fig. 2A). Somewhat surprisingly, the refined structure also revealed extension of the primer by dC (Fig. 3), indicating that hpol η possesses activity with Ca2+ as the metal ion cofactor. The primer strand makes a sharp turn at the active site to prevent a clash with the incoming dCTP, and the additional cytidine dC9 is located in the minor groove, where its base portion forms a H-bonding interaction with Arg-111 and stacks against the 2’-deoxyribose of template G4 (O6-MeG is residue 4; Fig. 3B).

In this insertion complex, O6-MeG and dCTP adopt a wobble-like, slightly sheared configuration whereby the O6-Me

### Table 1

| Oligonucleotide name | Sequence |
|----------------------|----------|
| Primer               | 5’-GGG GCT CTT AAG CGT CAT-3’ |
| Template             | 3’-GCC CGA GCA TTC GCA GTA XTA CT-5’ |

**Oligonucleotides used for crystallographic studies**

| Structure name       | DNA sequence | Incoming nucleotide |
|----------------------|--------------|---------------------|
| O6-MeG:dCTP Insertion| 3’-TCG CAG TAC TAC -5’ | dCTP |
| O6-MeG:dTTP Insertion| 3’-TCG CAG TAC TAC -5’ | dTMP-NPP |
| O6-MeG:dC Extension  | 3’-TCG CAG TXG TAC -5’ | dCMP-NPP |
| O6-MeG:dT Extension  | 3’-TCG CAG TXG TAC -5’ | dCMP-NPP |

**Oligonucleotides used for kinetic studies**

| Oligonucleotide name | Sequence |
|----------------------|----------|
| Primer               | 5’-GGG GCT CTT AAG CGT CAT-3’ |
| Template             | 3’-GCC CGA GCA TTC GCA GTA XTA CT-5’ |

**Incorporation Opposite O6-Methyl G by Pol η**

- **Compositions**
  - We determined two structures of complexes between hpol η and 12-mer template-8-mer primer duplexes (see Table 1 for sequences) containing O6-MeG, with nucleoside triphosphates paired opposite the adducted residue. The atomic coordinates and structure factors have been deposited in the Protein Data Bank under codes 5L1I (insertion O6-MeG opposite dCTP), 5L1J (insertion O6-MeG opposite dTMP-NPP), 5L1K (extension after O6-MeG opposite dC), and 5L1L (extension after O6-MeG opposite dT). In the first, determined to a resolution of 2.79 Å, the incoming nucleotide is dCTP, and in the second, determined to 1.94 Å resolution, the incoming nucleotide is dTMP-NPP. A summary of crystal data, data collection, and refinement parameters is provided in Table 3, and examples of the quality of the final electron density in the active site region are shown in Fig. 2, A and B. Crystals of the complex with the non-hydrolyzable dTTP analog dTMP-NPP were obtained readily, and electron density maps generated after molecular replacement and initial rigid body refinement revealed a fully occupied nucleoside triphosphate molecule. Conversely, dCMP-NPP in combination with Mg2+ in the crystal of the other insertion-stage complex was only partially ordered, with electron density visible for the triphosphate moiety and part of the sugar but not for the base. Subsequently, we resorted to dCTP at higher concentrations in the presence of Ca2+ (to prevent nucleotide insertion) to obtain crystals of the ternary complex in which the entire incoming nucleotide was surrounded by electron density (Fig. 2A). Somewhat surprisingly, the refined structure also revealed extension of the primer by dC (Fig. 3), indicating that hpol η possesses activity with Ca2+ as the metal ion cofactor. The primer strand makes a sharp turn at the active site to prevent a clash with the incoming dCTP, and the additional cytidine dC9 is located in the minor groove, where its base portion forms a H-bonding interaction with Arg-111 and stacks against the 2’-deoxyribose of template G4 (O6-MeG is residue 4; Fig. 3B).

In this insertion complex, O6-MeG and dCTP adopt a wobble-like, slightly sheared configuration whereby the O6-Me
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FIGURE 1. LC-MS analysis of primer extension products. A–D, reconstructed chromatograms for the indicated ion. E–H, fragmentation patterns for each of the indicated primer extension products.

TABLE 2
LC/MS analysis of full-length extension opposite O<sup>6</sup>-MeG

| Sequence | Charge/theoretical m/z | Measured m/z | Peak area (most abundant ion) | % Total |
|----------|------------------------|--------------|-------------------------------|---------|
| TCA TGA  | −2/934.60              | 934.36       | 17,630                        | 47 (C)  |
|          | −3/622.73              | 622.91       |                               |         |
| TCA TGAA | −2/1091.21             | 1090.82      | 3,497                         |         |
|          | −3/727.14              | 727.00       |                               |         |
| TTA TGA  | −2/942.11              | 941.82       | 19,807                        | 53 (T)  |
|          | −3/627.74              | 627.64       |                               |         |
| TTA TGA A| −2/1098.72             | 1098.36      | 4,440                         |         |
|          | −3/732.14              | 732.27       |                               |         |
moiety is shifted into the major groove and O2 of C is shifted into the minor groove (Fig. 3A). This relative orientation of the two nucleobases allows for formation of two bifurcated H-bonds. In the first, N2(H) of O6-MeG donates to both the O1 and N3 acceptors of dCTP (distances of 2.9 Å and 3.2 Å, respectively). In the second, N4(H)2 of dCTP donates to both the N1 and N3 acceptors of dCTP (distances of 2.9 Å and 3.2 Å, respectively). Thus, the two insertion-stage complexes demonstrate that the active site of hpol η is capable of accommodating a range of chemistries and pairing configurations and that key residues such as Arg-61 exhibit considerable conformational freedom.

Crystal Structures of Extension-stage Ternary hpol η Complexes—The crystal structures of two extension-stage complexes with O6-MeG opposite dC and dT and followed by a dG:dCMP-NPP pair were determined at resolutions of 1.82 Å and 1.62 Å, respectively. Selected crystallographic data are summarized in Table 3, and examples of the quality of the electron density around the final models are depicted in Fig. 2, C and D. In the complex with cytidine, O6-MeG and dC display more shearing than in the dCTP insertion complex. This precludes formation of bifurcated H-bonds, and the O6-MeG:dCdC pair is stabilized by two H-bonds that link N1(H) (O6-MeG) and N4(dC) and N2(H) (O6-MeG) and N3(dC) (Fig. 5). By comparison, the pairing mode between O6-MeG and dT in the other extension complex is virtually identical to that seen in the insertion-stage structure with the adduct paired opposite dTMP-NPP (Figs. 3 and 6). The base pair at the −1 position is stabilized by two H-bonds between N1 (O6-MeG) and N3(dT) (dTMP-NPP) and the other between N2(H) (O6-MeG) and O2(dT) (dT). In both complexes, Arg-61 assumes an orientation that is more commonly observed in complexes of hpol η, namely a curled conformation that allows an offset stacking interaction with the base of the incoming nucleotide and salt bridge formation with its α-phosphate group.

### Table 3

| Data Collection | O6-MeG:dCTP insertion | O6-MeG:dTMP-NPP insertion | O6-MeG:dC extension | O6-MeG:dT extension |
|-----------------|------------------------|---------------------------|---------------------|---------------------|
| Wavelength (Å)  | 0.97856                | 1.07810                   | 0.97872             | 0.97872             |
| Space group     | P61,1                   | P61,1                      | P61,1               | P61,1               |
| Resolution (Å)  | 50-2.79 (2.79-2.84)     | 50-1.94 (1.94-1.97)        | 50-1.82 (1.82-1.85) | 50-1.62 (1.62-1.65) |
| Unit cell a (Å) | 99.09, 81.89            | 99.00, 81.70               | 98.46, 82.11        | 98.36, 82.03        |
| Number of atoms | 3,372/409               | 3,450/395                 | 3,412/390           | 3,433/391           |
| Data Collection |                        |                           |                     |                     |
| R-work          | 0.159 (0.226)           | 0.159 (0.183)             | 0.153 (0.203)       | 0.156 (0.187)       |
| R-free          | 0.228 (0.266)           | 0.201 (0.281)             | 0.187 (0.232)       | 0.193 (0.240)       |
| Number of atoms | 3,122/390               | 2,9/2 (Ca2+)              | 28/2                | 28/2                |
| Protein residues| 138/2                   | 410/1                     | 491/1               | 526/3               |
| R-factor        | 43.6                    | 430                       | 430                 | 430                 |
| Average         | 49.1                    | 33.6                      | 23.8                | 4.4                 |
| Protein/DNA     | 49.4/55.5               | 33.1/37.3                 | 22.9/30.9           | 23.1/33.3           |
| Allomorph (Å)   | 30.5/38.9               | 30.3/26.8                 | 11.2/8.5            | 13.3/10.9           |
| Water/glycerol  | 37.2/50.7               | 41.1/25.9                 | 32.0/15.0           | 34.6/33.3           |
| ReLU mean square | 0.010                   | 0.018                     | 0.018               | 0.017               |
| Bonds (Å)       | 1.5                     | 1.8                       | 1.8                 | 1.8                 |
| Angles (degrees)| 95                      | 97                        | 97                  | 98                  |
| Allowed (%)     | 4.8                     | 2.5                       | 2.8                 | 1.8                 |
| Outliers (%)    | 0.2                     | 0.5                       | 0.2                 | 0.2                 |
| PDB ID code     | 5L11                    | 5L11                      | 5L1K                | 5L1L                |

* Statistics for the highest resolution shell are shown in parentheses.
Kinetic analysis of hpol η catalyzed polymerization past O\(^{6}\)-MeG using either the catalytic core construct (present work) or full-length enzyme (16) shows efficient bypass synthesis and equal preference for either correct C or incorrect T. The roughly equal ratio of C to T incorporation sets hpol η apart from the Y-family pols Dpo4 from *S. solfataricus* and hpol η/H9259 that both display a clear preference for error-prone insertion of T over C opposite O\(^{6}\)-MeG (4-fold (9) and 6-fold (15), respectively). The preference for T by hpol η was rationalized by the Hoogsteen pairing mode of the nascent base pair at the active site of the enzyme. Thus, the O\(^{6}\)-MeG:dTTP pair is stabilized by a H-bond between N7 of the adduct and N3-H of thymine (15). By comparison, N7 of O\(^{6}\)-MeG is positioned opposite an acceptor atom (N3) in the pair with dCTP, although one can invoke a single H-bond upon protonation of C at lower values of pH.

The different preferences for C and T found for insertion opposite O\(^{6}\)-MeG with *S. solfataricus* Dpo4 and hpol η are somewhat surprising given similarities between the two enzymes in the bypass of many adducts, including cyclic pyrimi-
idine dimers (17, 18) and 8-oxoG (19, 20). However, as the analyses of the efficient and correct bypass of the 8-oxoG adduct by both pols demonstrated (19-fold (19) and 4-fold preference (19) for dCTP over dATP insertion opposite the adduct by Dpo4 and hpol/H9257, respectively), the mechanisms underlying the mostly error-free bypass can differ significantly. Thus, Dpo4 relies on the Arg-332 residue from the little finger domain to maintain 8-oxoG in the template strand in the anti conformation to allow formation of a canonical Watson-Crick base-pairing with incoming dCTP (19). Conversely, hpol/H9257 focuses less on the template residue but recruits Arg-61 from the finger domain to guide the position of the incoming nucleoside triphosphate by contacting either base moiety or \(-\)phosphate, suggesting an inherent conformational flexibility of the side chain in response to a particular adduct (19) (Fig. 7).

Looking at the base pairing modes of the \(\text{O}^6\)-MeG adduct with dCTP and dTMP-NPP as well as with dC and dT at the active site of hpol (Figs. 3–6), we note that each pair is stabilized by two H-bonds, consistent with similar rates and extents of incorporation by this pol (16). The adduct pairing modes of this adduct (11) with dCTP and dC differ somewhat in that the former pair exhibits bifurcated H-bonds with less shearing and the latter two individual H-bonds with more shearing (similar to a wobble pair, e.g. G:U in RNA, but with the O2 atom of dT jutting into the minor groove rather than N2 of G; Fig. 5B). In fact, the two pairing modes of \(\text{O}^6\)-MeG with dCTP and dC observed at the hpol \(\eta\) active site, with varying degrees of shearing, closely resemble those of \(\text{O}^6\)-EtG:dC pairs in the structure of a DNA dodecamer with Hoechst 33258 bound in the minor groove (11) (PDB ID code 128D). In that duplex one of the pairs displays the two bifurcated H-bonds (dCTP insertion complex; Fig. 3), and the other one is stabilized by two individual H-bonds, with more shearing between \(\text{O}^6\)-EtG and dC (dC extension complex; Fig. 5).

**FIGURE 3.** Sheared configuration of incoming dCTP and \(\text{O}^6\)-MeG in a hpol \(\eta\) insertion stage complex. A, the active site viewed into the DNA major groove. B, rotated by \(-180^\circ\) around the vertical axis and viewed into the minor groove. Selected active site residues are colored by atom with carbon atoms shown in purple (\(\text{O}^6\)-MeG), cyan (dCTP), or green (Arg-61 and Gln-38 from the finger domain as well as Asp/Glu coordinated to Ca\(^{2+}\) (brown spheres)). The remaining nucleotides are shown in yellow, except for dC9 (blue), which was added to the primer during crystallization. H-bonds involving the adducted residue are depicted as dashed lines.

**FIGURE 4.** Watson-Crick-like configuration of incoming dTMP-NPP and \(\text{O}^6\)-MeG in a hpol \(\eta\) insertion stage complex. A, the active site viewed into the DNA major groove. B, rotated by \(-90^\circ\) around the horizontal axis and viewed perpendicular to the base planes of dTMP-NPP and \(\text{O}^6\)-MeG. Selected active site residues are colored by atom with carbon atoms shown in purple (\(\text{O}^6\)-MeG), cyan (dTMP-NPP), or green (Arg-61 and Gln-38 from the finger domain as well as Asp/Glu coordinated to Mg\(^{2+}\) (pink spheres)). The remaining nucleotides are shown in yellow, and H-bonds involving the adducted residue are depicted as dashed lines.
In the crystal structures of extension-stage Dpo4 complexes with a template strand containing O6-MeG and paired to either dC or dT, the former shows two H-bonds with shearing (9), just like the O6-MeG:dC pair at the active site of hPol H9257 (Fig. 5). However, in the structure with dT the weak electron density around the thymine moiety indicated enhanced flexibility of the incoming nucleotide and perhaps reduced stability of the nascent base pair compared with O6-MeG:dC. This scenario is consistent with the preference of Dpo4 for dCTP relative to dTTP.

Although both Dpo4 complexes were trapped at the extension stage in the crystal structures, the ceiling of the active site in the pol from S. solfataricus lacks the two amino acids that protrude from the finger domain and contact the nascent base pair in the case of hPol H9257 (Gln-38 and Arg-61, e.g. Fig. 3). Gln-38 interacts with sugar O4' of the adduct in the insertion complexes (Figs. 3 and 4), and Arg-61 contacts the incoming nucleotide. However, the interaction modes observed for Arg-61 vis-à-vis dCTP and dTMP-NPP differ distinctly. In the former complex, the side chain is rotated toward the triphosphate moiety, and the guanidino group forms a salt bridge with the α-phosphate of dCTP (Fig. 3). In the complex with dTMP-NPP, the side chain adopts two orientations whereby the guanidino group in both is facing away from the triphosphate moiety (Fig. 4). In one orientation, the positive charge of the arginine side chain hovers directly above the space between the thymine and O6-methylguanine bases, in close vicinity of the O4 and O6 atoms (not H-bonded) and N3(H) and N1 atoms (H-bonded) (Fig. 4B). The particular position that the guanidino moiety assumes in the complex with dTMP-NPP is unique among structures of ternary hPol H9257 complexes (Fig. 7, #5) and may help relieve a potential repulsion between the O6 and O4 acceptors from O6-MeG and dTMP-NPP, respectively, in the major groove.

Arg-61, which protrudes from the active site ceiling, can swivel around and adopt a range of orientations that allows it to interact with the α-phosphate of the incoming dNTP and par-
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FIGURE 7. Conformational flexibility of finger residue Arg-61 at the active site of ternary hpol η-DNA-dNTP complexes. View into the major groove of the DNA template (left) primer (right) duplex with dNTP visible to the right of Arg-61 side chains that are drawn in stick mode. Two divalent metal ions (Mg²⁺ or Ca²⁺) between the 3'-terminal nucleotide of the primer strand and the triphosphate moiety of the incoming nucleotide are located at the center of crossing dashed lines that mark their coordination geometry. A total of 23 crystal structures with the following PDB ID codes were included in the overlay: 3Z1J, 5K1J, and 5L1L (O⁶-MeG, this work), 4DLF, 4DLG, 5DQ7, 5DQ8, 5DQ9, and 5DQ1 (O⁶-methyl- and O⁶-ethyl-T) (22), 24RM, 4RNN, and 4RNO (abasic site) (21), 4RU9 (Mεpapy-dG) (23), 4O3N (native G:dCMP-NPP Mg²⁺ complex), 4O30, 4O3P, 4O3Q, 4O3R, and 4O3S (8-oxoG) (17), 5D7G, 5D7G, and 5D9G (1,N⁴-etheno-da) (24), and 3MR3 (cys-syn-cyclobutane thymine dimer) (18). Arg-61 adopts various orientations that we have marked with numbers 1 (curled conformation and interacting with the α- and β-phosphate groups of the incoming 2‘-deoxynucleotide triphosphate, the most common position), 2 (curled conformation and stacked in an offset manner on the nucleobase of the incoming dNTP), 3 (extended conformation and interacting with the major groove edge of the template base at the 0 position, e.g. 8-oxoG (magenta carbons)), 4 (curled conformation and interacting with the template base at the +1 position, e.g. the second T of cys-syn-cyclobutane thymine dimer (orange carbons)), and 5 (curled conformation and fully stacked on the nucleobase of the incoming dNTP, e.g. O⁶-MeG and dTMP-NPP, this work (lilac carbons)).

Itally stack on its base moiety (the most common orientation) but also stretch and interact with the major groove edges of template and incoming base (e.g. 8-oxoG and dGTP; Ref. 17), reach over to the template base of a cyclic thymine dimer (18), or even form a salt bridge with a phosphate from the template strand (e.g. abasic site; Ref. 21) (Fig. 7). However, even taking into account this conformational freedom of Arg-61, the orientation of the guanidino moiety in the complex with dTMP-NPP allows a face-to-face stacking interaction with thymine pro- trudes. It contrasts with the situation in the dCTP complex of hpol η and the corresponding complex of Dpo4 with dTTP and may account for the more favorable incorporation of dT by the human pol compared with that from S. solfataricus.

Although the dCTP and dTMP-NPP complexes of hPol η were crystallized under somewhat different conditions (Ca²⁺ versus Mg²⁺, respectively, and varying concentrations of the nucleoside triphosphate), the positions of Arg-61 in the respective complexes may reflect the particular electronic features of the nascent O⁶-MeG:dC and O⁶-MeG:dT pairs. In any case, the extension in situ of the primer in the presence of dCTP and Ca²⁺ demonstrates that the conditions used to crystallize the complex did not preclude activity (Fig. 3). Primer extension under conditions that employed Ca²⁺ in place of Mg²⁺ were also observed for hpol η crystallizations with a template containing O⁴-EtT (22) and had previously been seen in the struc-
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were obtained from Jena Bioscience (Jena, Germany). [γ-32P] ATP (specific activity 3000 Ci/mmol) was purchased from PerkinElmer Life Sciences. Biospin columns were from Bio-Rad.

Assays—steady-state kinetic assays were done using varying concentrations of dCTP and dTTP as described in detail elsewhere (31, 32) for the estimation of kinetic parameters ($k_{cat}/K_m$), data points were fit to hyperbolic plots using non-linear regression in Prism (GraphPad, La Jolla, CA). The LC-MS analysis of extended primers was performed as described in detail elsewhere (33–35).

Crystallization—The hanging drop vapor diffusion technique was used to grow all crystals. Two dodecamer DNA sequences modified with the O6-MeG lesion either at the fourth or fifth position of the template and complementary primer strands were used in the crystallization experiments and are listed in Table 1. DNA template and primer oligonucleotide solutions were mixed in a 1:1 molar ratio in the presence of 10 mM sodium HEPES buffer (pH 7.5), 0.1 mM EDTA, and 50 mM NaCl and annealed at 85 °C for 10 min followed by slow cooling to room temperature. The DNA solution was then mixed with hpol η protein in a 1.2:1 molar ratio in the presence of 50 mM Tris-HCl (pH 7.5) containing 450 mM KCl and 3 mM DTT followed by the addition of either 5 μL of 100 mM MgCl₂ or 5 μL of 100 mM CaCl₂. The protein-DNA mixture solution was placed in an Amicon filter (Millipore, Billerica, MA), and the solution was concentrated to a final concentration of ~2 mg of protein/ml. Either dCTP or one of the non-hydrolyzable nucleoside triphosphate analogs was added to the concentrated mixtures containing Ca²⁺ or Mg²⁺. The ternary complex solution was mixed with an equal volume of reservoir solution containing 0.10 M sodium MES (pH 5.5), 5 mM CaCl₂ (or 5 mM MgCl₂), and 14–18% (v/v) PEG 2000 monomethyl ether and equilibrated against 500 μL of reservoir solutions. Crystals commonly appeared after overnight incubation at 18 °C and were allowed to grow for 1 week to several weeks. They were transferred to 25% glycerol (v/v) and then frozen in liquid nitrogen for data collection.

Data Collection and Structure Refinement—X-ray diffraction data were collected at 100 K on the 21-ID-D, 21-ID-F, or the 21-ID-G beam lines at the Advanced Photon Source (APS), Argonne National Laboratory (Argonne, IL). Data were processed using HKL2000 (36), and all data collection statistics are summarized in Table 3. Structures were determined by molecular replacement in MOLREP (37, 38) using the coordinates of the complex between hpol η and native DNA (PDB ID code 4O3N) as the search model. Structures were refined either with PHENIX (39) or REFMAC (40), and model building was carried out in COOT (41). Model statistics and geometric parameters are summarized in Table 3. All structural figures were generated with the program UCSF Chimera (42).

Author Contributions—A. P. crystallized the complexes and solved the structures, Q. Z. performed the kinetic and LC-MS analyses, and F. G. and M. E. conceived the studies and wrote the manuscript along with A. P.

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