

I Am Chem Soc. Author manuscript; available in PMC 2009 May 5.

Published in final edited form as:

J Am Chem Soc. 2007 December 19; 129(50): 15412-15413. doi:10.1021/ja072127v.

(R)-β,γ-Fluoromethylene-dGTP-DNA Ternary Complex with DNA Polymerase β

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DNA polymerases are crucial for maintaining the fidelity of genetic information encoded into DNA, and failure to repair aberrant bases in damaged DNA strands is notoriously implicated in oncogenesis. During base excision repair (BER), 1,2 DNA polymerase β (pol β) inserts the correct deoxynucleoside triphosphate (dNTP), replacing an excised damaged or mismatched nucleoside residue, with release of pyrophosphate. The smallest eukaryotic cellular DNA polymerase, pol β is an error-prone enzyme that is tightly down-regulated in normal somatic cells, but often overexpressed in human tumors, and has been the subject of extensive studies examining its roles in BER 2 and cancer. 3

Probes of molecular interactions with nucleic acid polymerases can be created by designed modifications in the structures of natural dNTPs. Although such analogues may be usefully modified in their nucleoside moieties to create active site probes, as shown for example by the recent work of Kool, 4 changes to the triphosphate group are of particular interest because this group is the locus of chemical transformation catalyzed in the polymerase active site. Replacement of the $P\alpha$ -O-P β bridging oxygen by a carbon atom (CXY) will prevent hydrolysis, whereas a $P\beta$ -CXY-P γ modification will alter the leaving group properties, depending on the nature of substituents X and Y, while conferring resistance 5 to dephosphorylation. The introduction of these substituents potentially may also enable entirely new bonding (or repulsive) active site interactions, not present with the natural nucleoside triphosphate, and thus could inform inhibitor design seeking to exploit pol β as a drug target.

Recently, a series of unfluorinated (X,Y=H,1) and fluorinated (X,Y=F,2;X,Y=H,F,3) (R) or F,H 4 (S)) β , γ -methylenebisphosphonate analogues of dGTP (Chart 1) and related compounds were used to examine leaving group effects on pol β catalysis and fidelity. These analogues are substrates of the polymerase, but release an X,Y-methylenebisphosphonate in place of the natural pyrophosphate leaving group. Several β , γ -CXY dNTP analogues were previously investigated as inhibitors of different DNA polymerases and viral reverse transcriptases, but the structures of the putative complexes formed during turnover were not determined. In previous studies, the CHX analogues (or more generally, CXY analogues where $X \neq Y$) have been used as the diastereomer mixtures always obtained synthetically when the

Supporting Information Available: Synthesis and characterization data for 1-7; crystallographic data for the complex of 3 with DNA pol β (PDB ID, 2PXI); computer docking results. This material is available free of charge via the Internet at http://pubs.acs.org.

Note Added in Proof: A PDB search for ligand C-F interactions with the guanidinium group of Arg has documented a number of examples underscoring the fluorophilic character of the Arg side chain: Müller, K.; Faeh, C.; Diederich, F. Science 2007, 317, 1881-1886.

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substituted methylenebisphosphonate is coupled to a dNMP, and their potential for a stereospecific interaction with the polymerase active site has received little or no attention.

In this Communication, we report X-ray crystallographic studies and computer docking simulations of terminal ddNMP primer-template pol β complexes with dGTP β , γ -fluoromethylenebisphosphonate analogues **3** and **4**, leading to the first demonstration of *stereospecificity* in dNTP β , γ -CHX analogue-enzyme complex formation.

(*R/S*)-β,γ-CHF-dGTP (**3/4**) (and also the corresponding CH₂ and CF₂ analogues) were prepared by DCC-mediated conjugation of the morpholidate^{8,9} of dGMP (**5**) in anhydrous DMSO with the tributylammonium salt of the appropriate methylenebisphosphonic acid, followed by two-stage preparative HPLC to achieve a high degree of purity with respect to contaminating nucleotides. The (mono- and difluoromethylene)bisphosphonic acids^{10,11} (**6b,7b**) were prepared using the more convenient¹² Selectfluor (**8**) in place of perchloryl fluoride¹⁰ to synthesize the intermediate fluorinated esters (**6a,7a**)^{10,11} from tetraisopropyl methylenebisphosphonate. In consistency with an early study of the mixed (*R/S*)-β,γ-CHF analogues of ATP,¹³ at pH 10 or higher, **3** and **4** could be individually detected and quantified (1:1) by their overlapping but resolvable ¹⁹F NMR resonances, exhibited as a pair of ddd with δ-218.61 and -218.77 ppm.

For protein crystallography, human pol β was overexpressed in *E. coli* and purified as described previously. ¹⁴ The double-stranded DNA substrate consisted of a 16-mer template (5'-CCGACCGCGCATCAGC-3'), a complementary 9-mer primer (5'-GCTGATGCG-3'), and a 5-mer downstream oligonucleotide (5'-pGTCGG-3'), thus creating a two-nucleotide gap with annealed primer. Addition of ddCTP terminates the primer and creates a one-nucleotide gap. The 1:1 mixture of **3** and **4** was added to a solution of the preformed protein-DNA complex.

The crystal structure of the resulting complex was resolved at 2.1 Å. The analogue was found in the enzyme active site position normally occupied by dGTP, and its configuration overall was a good match for that of the natural substrate. However, despite both CHF stereoisomers being present at very similar concentrations in the crystallizing mixture, electron density was only observed for a single fluorine atom in the complex, corresponding to the (R)-CHF stereoisomer (3) (Figure 1). In contrast, in a comparable crystallography experiment, the corresponding β , γ -CHCl analogue stereoisomers populated the DNA pol active site about equally (Pedersen, L. C., personal communication). Exclusion of the (S)- β , γ -CHF analogue is not the result of an unattainable binding conformation, because the crystal structure of the corresponding CF₂-analogue complex superimposes well with that of the (R)-stereoisomer.

Thus, in the absence of an dominant steric factor, asymmetric polarization induced by the F substituent presumably influences **3** versus **4** binding specificity electrostatically. ¹⁵ If we assume that the limit for detection of fluorine electron density at the (S)-F position corresponds to a S/R ratio of roughly 1:4 or less, then a stereospecific interaction on the order of 0.8 kcal/mol would suffice. The fluorine atom in the **3** complex is located 3.1 Å from an Arg183 guanidine N atom (Figure 1), raising the possibility that an unusual ¹⁶ F•••H bonding interaction ¹⁷ contributes to stabilizing the preferred stereoisomer within the desolvated and preorganized ^{15,18} enzyme active site complex. Further studies are in progress to explore this possibility and alternative explanations. ^{16b}

Molecular docking calculations 19 are often valuable in studying protein-ligand interactions. Prior to attempting the crystallography studies, we first carried out an exploratory ligand docking experiment based on the α,β -NH-dUTP-DNA-pol β structure (2FMS), 20 using Autodock 3.0, 21 substituting β,γ -CF2-dTTP for the dUTP analogue. Docking runs predicted a preferred β,γ -CF2 triphosphate chain orientation similar to that of natural dNTPs, but placing one of the two diastereotopic F atoms close to Arg183 in the active site environment. Redocking

of 3 using our recently available β,γ -CF₂-dGTP-DNA-pol β structure (2ISO)⁶ revealed a clustering of solutions placing the F atom within bonding proximity of the Arg183, whereas with 4 such an interaction was less favored. An overlay of the nucleoside moieties and triphosphate backbones of 1, 2, and 3 within the DNA pol β complexes (X-ray crystallographic data) reveal them to be substantially congruent, confirming that introduction of the F atom(s) does not perturb the overall fit of the substrate to the active site and that the F atom positions are similar in 2 and 3.

In conclusion, under crystallization conditions, **3** is preferentially bound from a 1:1 mixture of diastereomers **3** and **4** into a DNA-pol β complex, in which a polar CHF bond to Arg183 is spatially allowed. Docking simulations predicted this configuration to be more likely with **3** than with its *S* stereoisomer **4**, which was not observed in the crystal complex. Substitution of a single fluorine atom at the bridging carbon atom of a β , γ -CH₂-dNTP analogue, while offering the advantage pK_a properties more closely mimicking those of the dNTP substrate, ²² also may result in stereospecific binding to the targeted active site, determined by the CHF chirality.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgment

Dedicated to Professor F. H. Westheimer, 1912-2007. We thank Dr. Kym Faull and Dr. Ron New for assistance with HRMS analysis. This research was supported by NIH Grant 5-U19-CA105010 and in part by the Intramural Research Program of the NIH, National Institute of Environmental Health Sciences.

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1: X,Y = H 3: X = H,Y = F

2: X,Y = F 4: X = F,Y = H

Chart 1.

β,γ-CXY Analogues of dGTP (Charges Omitted)

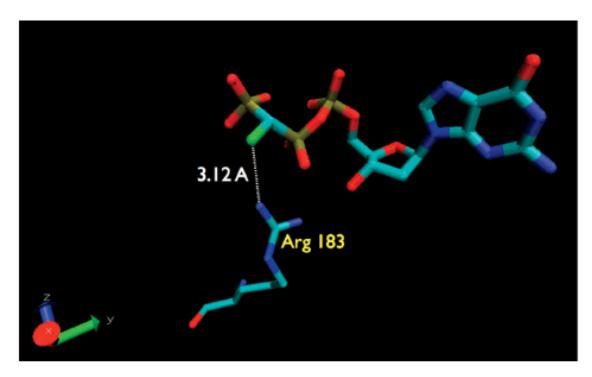


Figure 1. Detail from X-ray crystal active-site structure of DNA pol β :DNA complex soaked with a 1:1 mixture of 3 and 4.