Kinetics of deamination and Cu(II)/H$_2$O$_2$/Ascorbate-induced formation of 5-methylcytosine glycol at CpG sites in duplex DNA

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ABSTRACT

Mutation in p53 tumor suppressor gene is a hallmark of human cancers. Six major mutational hotspots in p53 contain methylated CpG (mCpG) sites, and C→T transition is the most common mutation at these sites. It was hypothesized that the formation of 5-methylcytosine glycol induced by reactive oxygen species, its spontaneous deamination to thymine glycol and the miscoding property of the latter may account, in part, for the ubiquitous C→T mutation at CpG site. Here, we assessed the kinetics of deamination for two diastereomers of 5-methylcytosine glycol in duplex DNA. Our results revealed that the half-lives for the deamination of the (5S,6S) and (5R,6R) diastereomers of 5-methylcytosine glycol in duplex DNA at 37°C were 37.4 ± 1.6 and 27.4 ± 1.0 h, respectively. The deamination rates were only slightly lower than those for the two diastereomers in mononucleosides. Next, we assessed the formation of 5-methyl-2'-deoxyxycytidine glycol in the form of its deaminated product, namely, thymidine glycol (Tg), in methyl-CpG-bearing duplex DNA treated with Cu(II)/H$_2$O$_2$/ascorbate. LC-MS/MS quantification results showed that the yield of Tg is similar as that of 5-(hydroxymethyl)-2'-deoxyxycytidine (Scheme 1). Together, our data support that the formation and deamination of 5-methylcytosine glycol may contribute significantly to the C→T transition mutation at mCpG dinucleotide site.

INTRODUCTION

Mutation in p53 tumor suppressor gene is common in human cancers, and the cytosines at CpG sites in the p53 gene are methylated. Although CpGs are underrepresented by 5-fold of their expected frequency in mammalian DNA, six major mutational hotspots in p53, i.e. codons 175, 213, 245, 248, 273 and 282, contain CpGs (1). The methylated CpGs (mCpG), therefore, are the most important mutation targets in p53.

Reactive oxygen species (ROS) constitute an important endogenous source of DNA damage (2), and an array of ROS-induced DNA lesions have been identified and characterized (3). The main species responsible for the ROS-induced DNA damage in cells appear to be hydroxyl radical generated by a Fenton-type reaction involving the reduction of H$_2$O$_2$ by transition metal ions (e.g. Fe$^{2+}$ and Cu$^{+}$) (4). In this context, ROS can damage 5-methylcytosine to give methyl group oxidation products and lead to the saturation of the C5 = C6 double bond, the latter gives 5-methyl-5,6-dihydroxy-5,6-dihydro-2'-deoxycytidine (or 5-methyl-2'-deoxyxycytidine glycol, mdCg; Scheme 1). The saturation of the C5 = C6 double bond in mdCg renders it susceptible to deamination, which affords 5,6-dihydroxy-5,6-dihydrothymidine (or thymidine glycol, Tg; Scheme 1) (5). Thymidine glycol is mostly a blocking lesion for replicative DNA polymerases (6). It, however, can be bypassed by translesion synthesis DNA polymerases including yeast DNA polymerases η and ζ as well as human DNA polymerase κ, and all three polymerases preferentially insert a dAMP opposite the lesion (7–9). Therefore, the formation and subsequent deamination of mdCg are thought to be one of the pathways leading to the C→T transition mutation at the CpG dinucleotide site. However, it remains unexplored how cytosine methylation affects the formation and deamination of the ring-saturated glycol derivative in duplex DNA.

Previous studies showed that the half-lives for the deamination of one diastereomer of 2'-deoxyxycytidine glycol and cytosine glycol (Cg) in calf thymus DNA at 37°C are 50 min and 28 h, respectively (10). On the other hand, the half-life for the deamination of the two diastereomers of mdCg in mononucleoside at 37°C is ~20 h (5). If the difference in deamination rate of Cg in mononucleoside and duplex DNA can be extrapolated to that...
for 5-methylcytosine glycol, then the deamination of 5-methylcytosine glycol in DNA may not contribute significantly to the C→T mutation at the CpG dinucleotide site. Therefore, understanding the implications of the deamination of 5-methylcytosine glycol in the ubiquitous C→T transition mutation at CpG sites necessitates the measurement of the rate of deamination of 5-methylcytosine glycol in duplex DNA.

Recently, we evaluated the formation of single-nucleobase lesions, i.e. 5-(hydroxymethyl)-2'-deoxycytidine (5-HmdC) and 5-formyl-2'-deoxycytidine (5-FodC), and tandem intrastrand cross-link lesions, i.e. G[8-5m]mC and mC[5m-8]G, emanating from the 5-methyl radical of mC at CpG sites (11). We found that the yield of single-nucleobase lesions is ~100 times more than intrastrand cross-links when methyl-CpG-bearing DNA was treated with Fenton-type reagents (11). A recent theoretical study from what was observed for Cg.

Scheme 1. ROS-induced DNA lesions of 5-methyl-2'-deoxycytidine and the deamination of 5-methyl-2'-deoxycytidine glycol.

**MATERIALS AND METHODS**

**Materials**

CuCl₂, L-methionine, L-ascorbic acid and nuclease P1 were from Sigma–Aldrich (St. Louis, MO, USA). Hydrogen peroxide (30%) was purchased from Thermo Fisher Scientific (Waltham, MA, USA). Common reagents for solid-phase DNA synthesis were obtained from Glen Research (Sterling, VA, USA). ODNs incorporated with an isolated Tg or Tg-(8-oxodG) tandem lesion were synthesized previously (16).

**Treatment of synthetic ODNs with Fenton-type reagents**

A self-complementary ODN, d(AXGTAXGTAXGTA XGT) (‘X’ represents 5-methylcytosine), was annealed in a buffer containing 50 mM NaCl and 20 mM phosphate (pH 6.9) by heating the solution to 80°C and cooling slowly to room temperature. Aliquots of ODNs (2.5 nmol) were incubated with CuCl₂ (6.25–100 μM), H₂O₂ (50–800 μM) and ascorbate (0.5–8 mM) in a 100 μl buffer (Table 1), which contained 50 mM NaCl and 20 mM phosphate (pH 6.9), at room temperature for 60 min. All chemicals were freshly dissolved in doubly distilled water and the reactions were carried out under aerobic conditions. The reaction was terminated by adding an excess amount of L-methionine, and the ODN samples were desalted by ethanol precipitation.

Table 1. Concentrations of Fenton-type reagents employed for the treatment of DNA

<table>
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<th>Control</th>
<th>A</th>
<th>B</th>
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<td>Cu(II) (μM)</td>
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<tr>
<td>H₂O₂ (μM)</td>
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<tr>
<td>Ascorbate (mM)</td>
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<td>2.0</td>
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<td>8.0</td>
</tr>
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</table>

*All reactions were carried out in a 100 μl solution containing 25 μM ODN, 25 mM NaCl and 50 mM phosphate (pH 6.9).
Deamination of mdCg and enzymatic digestion

A 27-mer duplex ODN, 5'-ACGGGCCGAAACXGGAGGGCCCGACCGT-3'/5'-ACGGTGGCCGACTGCGGCTTTTGCCCGACCGT-3' (X represents 5-methylcytosine, 3 nmol), was treated with Cu(II)/H2O2/ascorbate as described above (condition E in Table 1). The resulting solution was exchanged into a 100 μl buffer solution containing 10 mM phosphate (pH 6.9) by using YM-10 Centricon membrane filters. The solution was subsequently incubated at 37°C. Aliquots were removed at specific time points, frozen immediately on dry ice and stored at −80°C. Prior to mass spectrometric analysis, 0.5 U of nuclease P1 and a 1.5 μl solution containing 300 mM sodium acetate (pH 5.0) and 10 mM zinc acetate were added to 25 μl of the ODN sample. The digestion was continued at room temperature for 40 min, and the enzymes were removed by passing through a YM-10 Centricon membrane filter (Millipore, Billerica, MA, USA).

Generation of authentic Tg-containing ODNs

The ODN d(ATGGCTgGGCTAT), which contained the (5R,6S) diastereomer of the Tg, was synthesized previously (16,17). ODNs containing the two cis diastereomers of Tg were obtained by OsO4 oxidation (18). Briefly, 50 nmol d(ATGC) was dissolved in a 100 μl aqueous solution of 2% OsO4 and incubated at room temperature for 24 h before adding excess amount of NaHSO3 to reduce the OsO4. The residual OsO4 was removed by three times of extraction with equal volumes of carbon tetrachloride, and the aqueous layer was dried by using a Savant SpeedVac (Savant Instruments Inc., Holbrook, NY, USA). The dried residue was desalted by passing through HPLC and digested with nuclease P1 (vide supra) for the subsequent LC-MS/MS analysis.

Synthesis of ODNs containing Tg and a 3' neighboring dG that is uniformly labeled with 15N (ODNX)

Uniformly 15N-labeled dGTP, i.e., [2-amino-1,3,7,9,15N5]deoxyguanosine 5'-triphosphate (Cambridge Isotope Laboratories, Andover, MA, USA), was enzymatically added to the 3'-terminus of a synthetic tetrameric ODN d(GCAT) using terminal deoxynucleotidyl transferase (New England Biolabs, Ipswich, MA, USA) (19). The extension products were then treated with OsO4 to afford the Tg-containing ODNs (vide supra). The enzyme was removed by passing through YM-10 Centricon membrane filter and the ODNs were recovered by ethanol precipitation, desalted on HPLC, and used as internal standard for LC-MS quantification.

LC-MS and MS/MS analysis of p(mdCg)pdG and pTgp(dG)

The digestion mixture was analyzed by LC-MS/MS with an Agilent 1100 capillary HPLC system and an LTQ linear ion-trap mass spectrometer (Thermo Fisher, San Jose, CA, USA) equipped with an electrospray ionization source. A 0.5 x 150 mm Zorbax SB-C18 column (5 μm in particle size, Agilent Technologies, Palo Alto, CA, USA) was used for the separation of the above nuclease P1 digestion mixtures, and the flow rate was 6.0 μl/min. For deamination analysis, a 5-min gradient of 0–15% methanol in 400 mM 1,1,1,3,3,3-hexafluoro-2-propanol (HFIP, pH was adjusted to 7.0 by the addition of triethylamine), followed by a 35-min linear gradient of 15–35% methanol in 400 mM HFIP, was employed for the separation. The mass spectrometer was operated in the negative-ion mode, and the instrument was set up to acquire MS in the high-resolution ‘ultra zoom scan’ mode in the m/z range of 680–688 and the tandem mass spectra (MS/MS) for the fragmentation of the [M − H] ions of p(mdCg)pdG and pTgp(dG). The temperature for the ion transport tube of the mass spectrometer was maintained at 300°C.

For quantitative analysis, the digestion mixtures, to which 4 μl of ODNX was added prior to nuclease P1 digestion, was incubated at 37°C for over 60 h to allow for the extensive deamination of p(mdCg)pdG to pTgpG. The mass spectrometer was set up for monitoring the fragmentation of the [M − H] ions of the stable isotope-labeled pTgpG* as well as unlabeled dinucleotides pTgpdG and pTgp(8-oxodG). The calibration curves were constructed by subjecting, to LC-MS/MS analysis, the same amount of the isotope-labeled ODNX solution (4 μl) and different amounts of the authentic dodecameric standard d(ATGGCTTGCTAT) or d(ATGGCTG5G5GCTAT) (‘G5’ represents 8-oxodG) containing single Tg or tandem Tg-(8-oxodG) lesion, respectively (16). A 5-min gradient of 2–20% methanol in 400 mM HFIP, followed by a 35-min gradient of 20–50% methanol in 400 mM HFIP, was employed for the separation.

RESULTS

Kinetics of deamination for the two cis diastereomers of mdCg in duplex ODNs

To measure the kinetics of deamination of 5-methylcytosine glycol in double-stranded DNA, we first produced the 5-methylcytosine glycol-carrying duplex DNA by treating 3 nmol of a 27-mer duplex ODN, 5'-ACGGGCCGAAACXGGAGGGCCCGACCGT-3'/3'-TGCCCGGCTTTGGXCTCCGGGCTGGCA-5', where ‘X’ is a 5-methylcytosine, with Fenton-type reagents (100 μM CuCl2, 800 μM H2O2 and 8 mM ascorbate) in a buffer at room temperature for 60 min. The reaction was quenched with methionine, and the solution was buffer-exchanged into a 10-mM phosphate buffer and incubated at 37°C to allow for the deamination to occur. Aliquots were removed at different time points, frozen immediately on dry ice and stored in a −80°C freezer. After a quick digestion with nuclease P1, samples were subjected immediately to on-line LC-MS analysis, where the dinucleotide p(mdCg)pdG and its deaminated counterpart, i.e., pTgpG, were monitored in a high-resolution ‘zoom-scan’ mode on an LTQ linear ion-trap mass spectrometer. In this respect, it was found that the bulky nature of thymidine glycol and its lack of aromaticity prohibit the nuclease P1-mediated cleavage of its 3‘ phosphodiester.
linkage (18–20); thus, the digestion with nuclease P1 produces a dinucleotide containing the lesion along with its flanking 3’ unmodified nucleotide. Indeed HPLC analysis of the nuclease P1 digestion mixture of a dodecameric ODN harboring a single Tg showed that ~89% of the Tg was released as a dinucleotide pTgpdG (Figure S1). It is of note that the digestion with both nuclease P1 and alkaline phosphatase has been employed by Bailey et al. (19) for the LC-MS/MS identification of Tg formed in cultured mouse fibroblast cells. We, however, observed that only a small fraction of Tg is liberated as a dinucleoside monophosphate after the lesion-carrying ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside, which might be attributed to the presence of residual endonuclease activity in the commercial preparation of the alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside. Indeed HPLC analysis of the ODN had been treated with both nuclease P1 and alkaline phosphatase; the majority of the lesion is released as a mononucleoside.

The tandem mass spectra of the [M – H]− ions of p(mdCg)pdG (m/z 683, (a)) and pTgpdG [m/z 684, (e)], as well as for monitoring the m/z 683 → 524, 567 transitions of p(mdCg)pdG (b) and for monitoring the m/z 684 → 541 transition of pTgpdG (d). The sample was the nuclease P1 digestion mixture of 27-mer mC-containing duplex DNA after it was treated with Fenton reagent and deamminated at 37 °C for 24 h (details are described in ‘Materials and Methods’ section). Shown also are the SICs for the m/z 684 → 541 transition for the analyses of the nuclease P1 digestion mixtures of an ODN, d(ATGCCGCTGAGCTAT), which contains the (5R,6S) diastereomer of Tg (e) and the nuclease P1 digestion mixture of d(ATGC) that has been treated with OsO4 (f). The MS/MS of the deprotonated ions of p(mdCg)pdG and pTgpdG are shown in Figures S2–S4.

Figure 1. The SICs for monitoring the [M – H]− ions of p(mdCg)pdG [m/z 683, (a)] and pTgpdG [m/z 684, (e)], as well as for monitoring the m/z 683 → 524, 567 transitions of p(mdCg)pdG (b) and for monitoring the m/z 684 → 541 transition of pTgpdG (d). The sample was the nuclease P1 digestion mixture of 27-mer mC-containing duplex DNA after it was treated with Fenton reagent and deamminated at 37 °C for 24 h (details are described in ‘Materials and Methods’ section). Shown also are the SICs for the m/z 684 → 541 transition for the analyses of the nuclease P1 digestion mixtures of an ODN, d(ATGCCGCTGAGCTAT), which contains the (5R,6S) diastereomer of Tg (e) and the nuclease P1 digestion mixture of d(ATGC) that has been treated with OsO4 (f). The MS/MS of the deprotonated ions of p(mdCg)pdG and pTgpdG are shown in Figures S2–S4.
Deamination was estimated to be described in ‘Materials and Methods’ section, and the initial extent of mer duplex sequence and deamination reaction conditions were averaged from the two major peaks found in Figure 1a and c. The 27-

...of the dinucleotide p(mdCg)pdG (m/z 0 432, arising from the neutral loss of a 2-deoxyadenosine, and an abundant fragment ion of m/z 330, being the [M – H] – ion of pdA. This normalization can facilitate the correction of difference in analyte loss during various steps of sample manipulation and the variation in instrument response. Plotting the normalized ratio versus the deamination time gives straight lines (Figure 3), supporting that the deamination follows a first-order reaction kinetics. The slope of the line facilitates the determination of the rate constant, which allows for the calculation of the half-life for the deamination of 5-methylcytosine glycol in duplex DNA (Figure 3 and the results are summarized in Table 2). It turned out that the rate constants for the deamination of the (5S,6S) and (5R,6R) diastereomers in duplex DNA at 37°C were 0.0186 and 0.0253 h⁻¹, respectively, and the half-lives for their deamination were calculated to be 37.4 and 27.4 h, respectively.

We also assessed the kinetics of deamination mdCg in duplex DNA by measuring the decrease in signal intensity for p(mdCg)pdG, and the rate constant for the deamination of the major and minor diastereomers of mdCg were found to be 0.0177 and 0.0147 h⁻¹, respectively (Figure S5). It remains unclear what accounts for the discrepancy in rate constants determined from the rise in signal for pTgpG and the drop in signal for p(mdCg)pdG. However, we speculate that this might be attributed, in part, to the presence of co-eluting isobaric impurities with p(mdCg)pdG and/or the lack of quantitatively release of mdCg from duplex DNA. We have shown that the nuclease P1-mediated release of pTgpG from ODN is nearly quantitative and the product-ion spectra of the pTgpG emanating from the deamination reaction are very similar as those of the standard compounds (vide supra). In addition, the increase in signal of pTgpG has to arise from the deamination of mdCg in duplex DNA. Therefore, we reason that the rate constants determined from the signal increase in pTgpG are reliable.

**LC-MS/MS quantification of pTgpG and pTgp(8-oxodG) lesions induced in duplex DNA by Cu(II)/H₂O₂/ascorbate**

To quantify the total amount of Tg formed from the deamination of mdCg, we treated a self-complementary duplex ODNs with different concentrations of Cu(II)/H₂O₂/ascorbate (Table 1), digested the resulting ODNs with nuclease P1 to release the lesions as dinucleotide pTgp(8-oxodG) lesions induced in duplex DNA. Plotted are the ratios of peak area found in the SIC for the m/z 684 – 541 transition (for pTgpG) over that found in the SIC for monitoring the fragmentation of the ion of m/z 330 (for pdA) versus the deamination time. The solid square and triangle represent the results based on the formation of pTgpG containing the (5S,6S) (the 19.5-min peak in Figure 1d) and (5S,6R) (the 20.1-min peak in Figure 1d) diastereomers of Tg (Scheme 1), respectively. The deamination reaction conditions were described in ‘Materials and Methods’ section.

Figure 2. High-resolution ‘zoom-scan’ ESI-MS for monitoring the kinetics of deamination of 5-methylcytosine glycol in a 27-mer duplex DNA. Shown are the regions of the mass spectra for the [M – H] – ions of the pTgpG (m/z 684) at 0 (a), 6 (b), 12 (c), 24 (d), 48 (e), and 72 h (f) after the initiation of the deamination reaction, and the spectra were averaged from the two major peaks found in Figure 1a and c. The 27-mer duplex sequence and deamination reaction conditions were described in ‘Materials and Methods’ section, and the initial extent of deamination was estimated to be ~20%.

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A fast chromatographic gradient was applied to allow for the two diastereomers to co-elute (Figure 4). The SIC for the m/z 684 → 346, 541, 689 → 351, 546, and 700 → 362, 557 transitions, which monitor the losses of thymidine glycol 5'-phosphate and the C₅H₅NO₄ component, respectively (Figure S6), showed a peak at 18.2 min, which supported the formation of the pTgp(8-oxodG) tandem lesion. It is worth noting that the elimination of C₅H₅NO₄ is a characteristic neutral loss from thymine glycol-containing dinucleotides and ODNs (Figure S6) (18). The quantification results showed that the pTgpG exhibited a linear dose-responsive increase when the concentration of H₂O₂ was up to 800 μM; however, the yield of pTgpG(8-oxodG) exhibits a nonlinear response over the dose range of Fenton-type reagents used (Figure 5, and calibration curves for LC-MS/MS quantification are shown in Figure S7). The yields of pTgpG(8-oxodG) at low-dose range increase slowly with the rise in the concentrations of Cu(II)/H₂O₂/ascorbate. However, when DNA was treated with the Fenton-type reagent at the three higher doses, the yield of pTgpG(8-oxodG) increased sharply. At the highest dose, the yield of pTgpG(8-oxodG) is ~10% of that of pTgpG. The exact reason for the disproportionally high yield at higher doses remains ambiguous, though it is possible that the formation of mdCg at the 5-methyl-2'-deoxycytidine site may promote the oxidation of its neighboring 2'-deoxyguanosine, and vice versa.

Hydroxyl radical can be added to C5 = C6 double bond of mC to form the glycol lesions, and it can also abstract a hydrogen atom from the methyl group of mC to generate an allyl radical. The allyl radical can result in the formation of two other single-nucleobase lesion, i.e. 5-HmdC.

**Table 2.** Kinetic parameters for the deamination of 5-methylcytosine and 5-methylcytosine glycol at 37°C

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<th>mC in Duplex DNAᵇ</th>
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<tr>
<td></td>
<td>(5S,6S)</td>
<td>(5R,6R)</td>
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ᵃData from ref. (5).
ᵇData from ref. (28).

**Figure 4.** SICs for monitoring the m/z 684 → 346, 541 [(a), for unlabeled pTgpG], m/z 689 → 351, 546 [(b), for labeled pTgpG] and m/z 700 → 362, 557 [(c), for unlabeled pTgpG(8-oxodG)] transitions obtained from the LC-MS/MS analyses of the nuclease P1 digestion mixture of Fenton-reagent-treated ODNs under reaction condition C (Table 1).

**Figure 5.** Cu(II)/H₂O₂/ascorbate-induced formation of pTgpG (a) and pTgpG(8-oxodG) (b) lesions in ODN d(AXGTAXGTAXGTAXGT) (‘X’ represents 5-methylcytosine). The values represent the means ± SD from three independent oxidation and quantification experiments.
and 5-FodC (Scheme 1), as well as two intrastrand cross-links, i.e. G[8-Sm]mC and mC[5m-S]G (11,23,24). To compare the damage products from different mC radicals, we also measured yields of 5-HmdC in the same ODN sequence. It turned out that the yield of Tg is comparable to that of 5-HmdC under the same experimental conditions, but was 5–6-fold lower than that of 8-oxodG (Figures 5a and Figure S8, and calibration curves are depicted in Figure S9). It is worth noting that the quantification results for 5-HmdC and 8-oxodG are consistent with the data that we obtained earlier for ODNs in different sequence contexts (11).

**DISCUSSION**

After the saturation of the C5 = C6 double bond, cytosine derivatives are susceptible to hydrolytic deamination. It was observed recently that the half-life of Cg is greatly enhanced in DNA, which is increased by 34-fold from 50 min in free nucleoside to 28 h in calf thymus DNA (10). A more recent study indicated that the half-life of Cg in duplex structure is still approximately eight times longer than Cg in free nucleoside. On the other hand, we observed that the deamination of the (S,6S) diastereomer of mdCg in duplex DNA is merely 2-fold slower than the same compound in mononucleoside, whereas the place-ment of the (S,6S) diastereomer in duplex DNA only increases the lifetime of the lesion by approximately 20% (Table 2). The striking difference in the effect of duplex structure on the deamination of Cg and 5-methylcytosine glycol suggests that the presence of a methyl group on the C5 carbon in 5-methylcytosine glycol, along with the saturation of the C5-C6 bond, may facilitate the lesion to adopt an extrahelical conformation in duplex DNA, thereby enabling the facile deamination of the lesion. Along this line, a previous NMR structural study demonstrated that the structurally related thymidine glycol is largely extrahelical (26). It is of note that, in isolated nucleoside, the (S,6S) diastereomer is deaminated at a slightly slower rate than the (S,6R) diastereomer, whereas the opposite is true when the two diastereomers are in duplex DNA. The difference might be due to the different steric hindrance experienced by the two diastereomers in duplex DNA, which may affect the nucleophilic attack of mdCg by a water molecule and its subsequent deamination to Tg. Future structural studies of duplex DNA housing the two diastereomers of mdCg or Tg will offer important insights about the difference of the two diastereomers of mdCg toward hydrolytic deamination.

As the presence of Tg inhibits the cleavage of its 3’ phosphodiester linkage by nuclease P1, which results in the liberation of the lesion as a dinucleotide bearing the 3’-flanking nucleotide (18-20), we quantified the Tg lesions formed at mCpG site as pTgpdG dinucleotide using negative-ion mass spectrometry. This also allows us to quantify the Tg-(8-oxodG) tandem lesion formed at mCpG site. In our study we used synthetic ODN containing an isotope-labeled 5’-pTgpdG*’-3’ sequence (‘DG’ represents the uniformly 15N-labeled 2’-deoxyguanosine) as internal standard, this method can effectively eliminate the error introduced by the potential lack of quantitative release of the lesion as a dinucleotide by nuclease P1 and spontaneous dephosphorylation during various stages of sample manipulation.

Previous experimental data obtained for poly(dG-mdC) upon exposure to γ rays (13) and theoretical calculations at the mononucleoside level (12) indicated that the hydroxyl radical can attack 5-methylcytosine at the C5 = C6 double bond and the 5-methyl group, with preference being on attack at the double bond to yield 5-methylcytosine glycol. The latter can deaminate to afford thymine glycol. However, the yield of thymine glycol in ODNs treated with Fenton-type reagents amounts to about five to six lesions per 1000 nucleosides at the highest dose, and is in a similar range as the other two single-nucleobase lesions arising from the 5-methyl radical of mC, i.e. 5-hydroxymethylcytosine (Figure 5a, Figure S8, and Table S1) and 5-formylcytosine (11). This result may indicate that the theoretical calculation at the mononucleoside level does not reveal completely the reactivity of the nucleoside in duplex DNA, where the methyl group in 5-methylcytosine, located in the major groove, can be more accessible to hydroxyl radical attack than the C5 = C6 double bond.

The efficient formation of mdCg and its facile deamination to Tg, together with the production of 5-HmdC and 5-FodC, may account for the prevalent C → T transition mutation found at CpG site in human p53 gene, a hallmark for many types of human tumors (27). In this context, several mechanisms have been proposed to account for the ubiquitously observed C → T transition mutation at CpG site (1). Two of them involve the deamination processes. One is the direct deamination of 5-methylcytosine to thymine. In support of this mechanism, Shen et al. (28) found that the rate for the deamination of 5-methylcytosine in duplex DNA at 37°C is 3.3×10⁸ h⁻¹, which is twice as fast as the corresponding deamination of cytosine. The other involves the formation of mdCg and its subsequent deamination to Tg, and we showed that this latter deamination is about 7 orders of magnitude faster than the spontaneous deamination of 5-methylcytosine in duplex DNA (Table 2). For the two deamination pathways to be equally important in contributing to the C → T transition mutation at CpG site, mdCg has to form at a frequency of one lesion per 10⁷ 5-methyl-2’-deoxycytidine. Our quantification results showed that the rate for the formation of mdCg is about 5–6-fold lower than that of 8-oxodG, and a recent careful study showed that the basal level of 8-oxodG in human cells is approximately 2.2 lesions per 10⁷ 2’-deoxyguanosine (29). Thus, the two deamination pathways may make a similar extent of contribution to the frequently observed C → T mutation at CpG site. It is worth noting that the above analysis is under the assumption that the product arising from the spontaneous deamination of 5-methylcytosine, i.e. a G:T mismatch, and the product generated from the deamination of mdCg, i.e. a G:Tg pair, are repaired.
in cells at a similar efficiency. Further studies are required to assess whether this is indeed the case.

It is also important to discuss how the presence of an additional methyl group on the C5 carbon affects the ROS-induced damage to cytosine residues in DNA. The results presented in this article, together with our recently published data (11), revealed that the total yield of the single-nucleobase lesions, Tg, 5-HmdC and 5-FodC, emanating from mCpg in duplex DNA upon exposure to Cu(II)/H2O2/ascorbate is several fold higher than the major single-nucleobase lesion, 5-hydroxy-2-deoxyuridine (5-OHdC), formed at unmethylated cytosine at CpG sites. A mutation frequency of 0.05% was reported for 5-OHdC after the lesion-bearing single-stranded M13 genome was propagated in Escherichia coli cells (30), and a mutation frequency of 0.03–0.28% was found for 5-FodC after the damage-harboring double-stranded shuttle vector was replicated in simian COS-7 cells (31).

The mutagenic properties of 5-HmdC and mdC have not yet been examined; however, the deaminated product of mdC (i.e. Tg) is known to direct the predominant incorporation of adenine nucleotide during DNA replication (7–9). Together, the enhanced formation of oxidatively generated lesion from methylated cytosine, combined with the facile deamination of mdC and the high mutagenic potential of the deaminated product, may render 5-methylcytosine more susceptible to produce C→T transition mutation than its unmethylated counterpart.

We also observed that the yield for the formation of the tandem lesion, i.e. Tg-(8-oxodG), is only about 10-fold less than that found for Tg that is 5’ neighboring to an unmodified dG (Figure 5). Therefore, the high yield for the formation of this tandem lesion at mCpg sites and the inefficient removal of 8-oxodG from Tg-(8-oxodG) site by human 8-oxoguanine DNA glycosylase (hOGG1) (32) may account for the high frequency of mCG→TT mutation observed after the CpG-methylated and Cu(II)/H2O2/ascorbate-treated pSP189 shuttle vector has been replicated in NER-deficient XPA cells (15). Along this line, both NER and long-patch BER repair pathway have been proposed for the repair of another type of tandem lesions composed of a thymine glycol and a neighboring 5'-2-deoxyribonolactone (33).

Copper plays a pivotal role in maintaining the structure and integrity of chromatin (34,35). Our results suggest that transition metal ion, copper or iron (11), can induce significant amount of DNA damage, thereby posing a great challenge for people suffering from Wilson’s disease and primary hemochromatosis, which cause abnormal accumulation of copper and iron in various organs (36,37). The tandem and/or cluster lesions, whose biochemical effects are largely unknown, constitute a major family of DNA damage products produced by Fenton-type reagents and ionizing radiation (38). The mutagenic properties of this particular type of tandem lesions are currently investigated in our laboratory.

Methylation pattern of cytosine residue at CpG sites is crucial for maintaining the epigenetic signal in human cells (39). The ROS-induced nucleobase modification at CpG sequence may also result in disturbed gene regulation and heritable epigenetic changes in chromatin (40).

**SUPPLEMENTARY DATA**

Supplementary Data are available at NAR Online.

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