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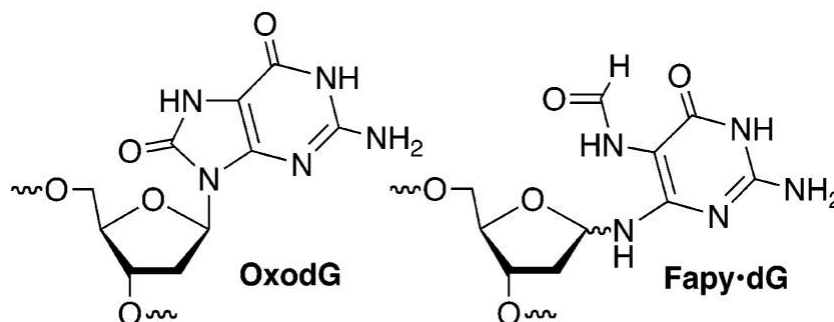
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Facile Quantification of Lesions Derived from 2'-Deoxyguanosine in DNA

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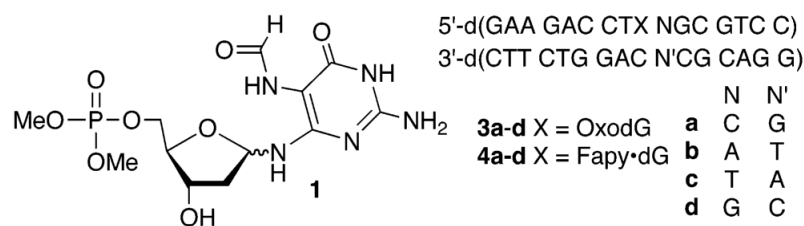
Exposure of DNA to free radicals and other electrophiles results in strand breaks, as well as modified nucleotides (lesions) within intact strands of the biopolymer. DNA lesions can be genotoxic and have been implicated in aging and diseases such as cancer.¹⁻³ OxodG and Fapy•dG are examples of lesions produced in significant yields from dG when DNA is exposed to oxidative stress. It is well-known that OxodG gives rise to G→T transversions.¹ Recent investigations suggest that Fapy•dG also has significant effects on repair and replication.⁴ For instance, OxodG and Fapy•dG result in high levels of G→T transversions when they are replicated in simian kidney cells.⁵ The former has been proposed as a biomarker when studying the effects of oxidative stress on DNA.² In addition, it has also been suggested that the ratio of OxodG:Fapy•dG may be characteristic of the cancerous state of a cell.⁶ Currently, these lesions are detected and quantified via mass spectrometry following degradation of the DNA.^{7,8} However, there is some disagreement regarding how these measurements are made.⁹⁻¹¹ We describe a reagent and accompanying fluorescent detection method that enables one to selectively quantify OxodG and Fapy•dG.



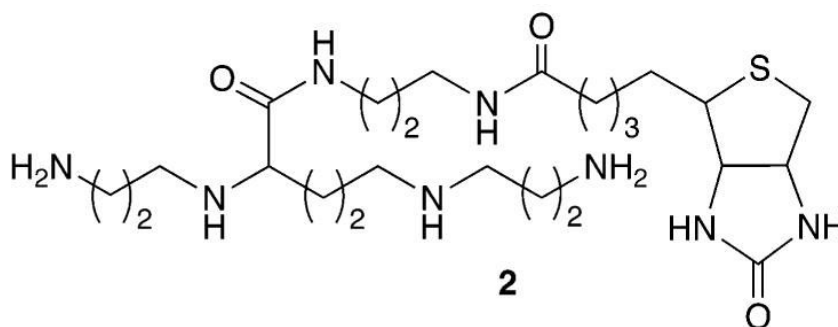
Selective DNA lesion detection is attractive because of its simplicity. One approach exploits a lesion's reactivity.¹²⁻¹⁴ The facile oxidation of OxodG and nucleophilic trapping of its oxidized product(s) by molecules such as spermine provided inspiration for developing a system for detecting it and Fapy•dG.¹⁵ OxodG (~0.75 V, NHE) is more readily oxidized than the most readily oxidized native nucleotide, dG (1.29 V, NHE). The irreversible oxidation of **1** (1.08 V, NHE) was determined to be more favorable than that of dG, suggesting that using a similar oxidation/trapping approach for detecting Fapy•dG was plausible.¹⁶

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 Supporting Information Available: Procedures for synthesis and characterization of all molecules, and all other experiments. ESI-MS and MALDI-TOF MS from experiments described herein. This material is available free of charge via the Internet at <http://pubs.acs.org>.



A carboxylated derivative of spermine was used as a precursor to **2**.^{16,17} The spermine derivative (**2**) provides a means for tagging the oxidized species derived from OxodG and Fapy•dG with biotin, which is used for quantifying the lesions. Slower moving adducts were detected in good yield when duplex DNA (**3** and **4**) was reacted with Na₂IrCl₆ in the presence of **2** at 25 °C for 1 h (Figures 1 and 2).¹⁵ Minor amounts of a product that migrated even more slowly were also detected. Reaction of OxodG with spermine and other diamines under oxidative conditions is well-established.¹⁵ Confirmation that adducts corresponded to 1:1 adducts between **2** and oxidized OxodG or Fapy•dG was obtained using MALDI-TOF MS.¹⁶ Additional evidence that adduct formation occurred at Fapy•dG in **4** was obtained by ESI-MS following reaction of **1** and **2** under the same reaction conditions.¹⁶



Although the yields of adducts were different for Fapy•dG and OxodG, for each lesion, they were independent of the flanking sequence (Figure 2). This was true even when OxodG was present in sequences that were expected to significantly alter its oxidation potential from that in **3a-d**.^{16,18} In addition, no adducts were detected when duplexes containing OxodA or Fapy•dA were reacted with the strongest oxidant, Na₂IrCl₆, in the presence of **2** (data not shown). Moreover, no adducts were observed when DNA containing 5-hydroxy-2'-deoxyuridine or 5-hydroxy-2'-deoxycytidine, which have more favorable oxidation potentials than OxodG, were reacted with Na₂IrCl₆ and **2** (data not shown).¹⁹ Their lack of reactivity under the reaction conditions indicates that oxidation is a necessary but insufficient requirement for tagging.

In order to quantify the individual amounts of Fapy•dG and OxodG, other oxidants were explored in the hope of taking advantage of the lesions' differing oxidation potentials. Although both produced adducts with **2** in the presence of Na₂IrBr₆, only OxodG was trapped when K₃Fe(CN)₆ was used as the oxidant (Figures 1 and 2). The selective biotinylation of DNA containing the lesions as a function of oxidant indicated that individual amounts of OxodG and Fapy•dG in randomly damaged DNA could be determined using **2** as part of a fluorescence assay (Scheme 1). Amplex Red, which is oxidized to fluorescent resorufin by horseradish peroxidase in the presence of H₂O₂, was used to quantify adducts of **2**.²⁰ Experiments were carried out immediately after tagging by **2** because the adducts decompose with half-lives on the order of 12–15 h.^{15,16} The system was calibrated for application in a microtiter plate using a 287 nt duplex fragment of M13mp7(L2) prepared by PCR that contained biotin at the 5'-

terminus of one strand.¹⁶ The fluorescent signal produced by resorufin depended linearly on the amount of biotinylated DNA as it varied 10-fold.

The ELISA type method was used to measure the amounts of OxodG and Fapy•dG produced in aqueous solution by γ -radiolysis. The amount of biotinylated DNA was measured as a function of dose using Na_2IrCl_6 and $\text{K}_3\text{Fe}(\text{CN})_6$ as oxidants (Figure 3). The adduct yield varied linearly with dose when either oxidant was used. The amounts of OxodG and Fapy•dG per dose (Gray) of radiation were determined (eqs 1 and 2) using the average yields of adduct formation established from experiments with **3** and **4** (Figure 2).¹⁶ Femtomoles of adduct(s) were readily detectable using this method. The data reveal that the ratio of Fapy•dG to OxodG formed in

$$\text{Yd. Adduct}(\text{Na}_2\text{IrCl}_6) = (0.88)\text{OxodG} + (0.81)\text{Fapy}\cdot\text{dG} \quad (1)$$

$$\text{Yd. Adduct}(\text{K}_3\text{Fe}(\text{CN})_6) = (0.79)\text{OxodG} \quad (2)$$

aqueous solution by ^{137}Cs equals 1.12. This is higher than the ratio reported when chromatin is irradiated but lower than when monocyte cells were subjected to γ -radiolysis.^{21,22} In summary, we have developed a simple and sensitive method for quantifying OxodG and Fapy•dG, two important DNA lesions. This tool will be widely accessible to researchers studying the effects of oxidative stress on DNA. In addition, the fluorescence assay should be generally applicable to the quantification of other lesions that can be selectively tagged with biotin.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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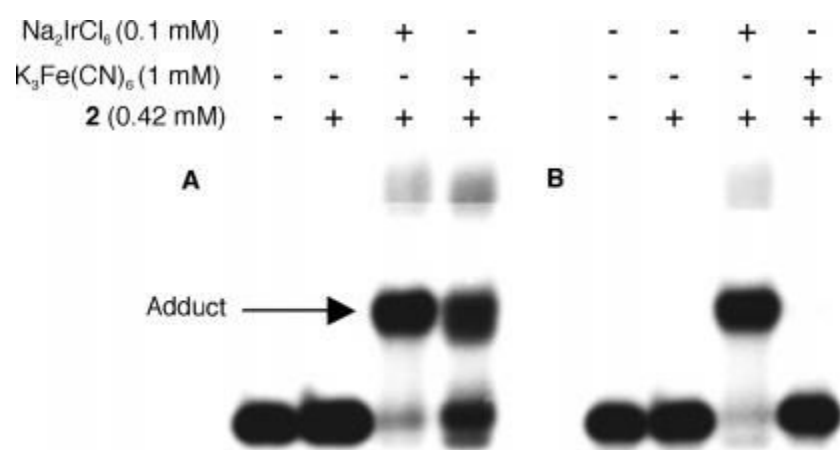


Figure 1. Reaction of **2** with DNA containing (A) OxodG (**3b**) or (B) Fapy•dG (**4b**) under oxidative conditions.

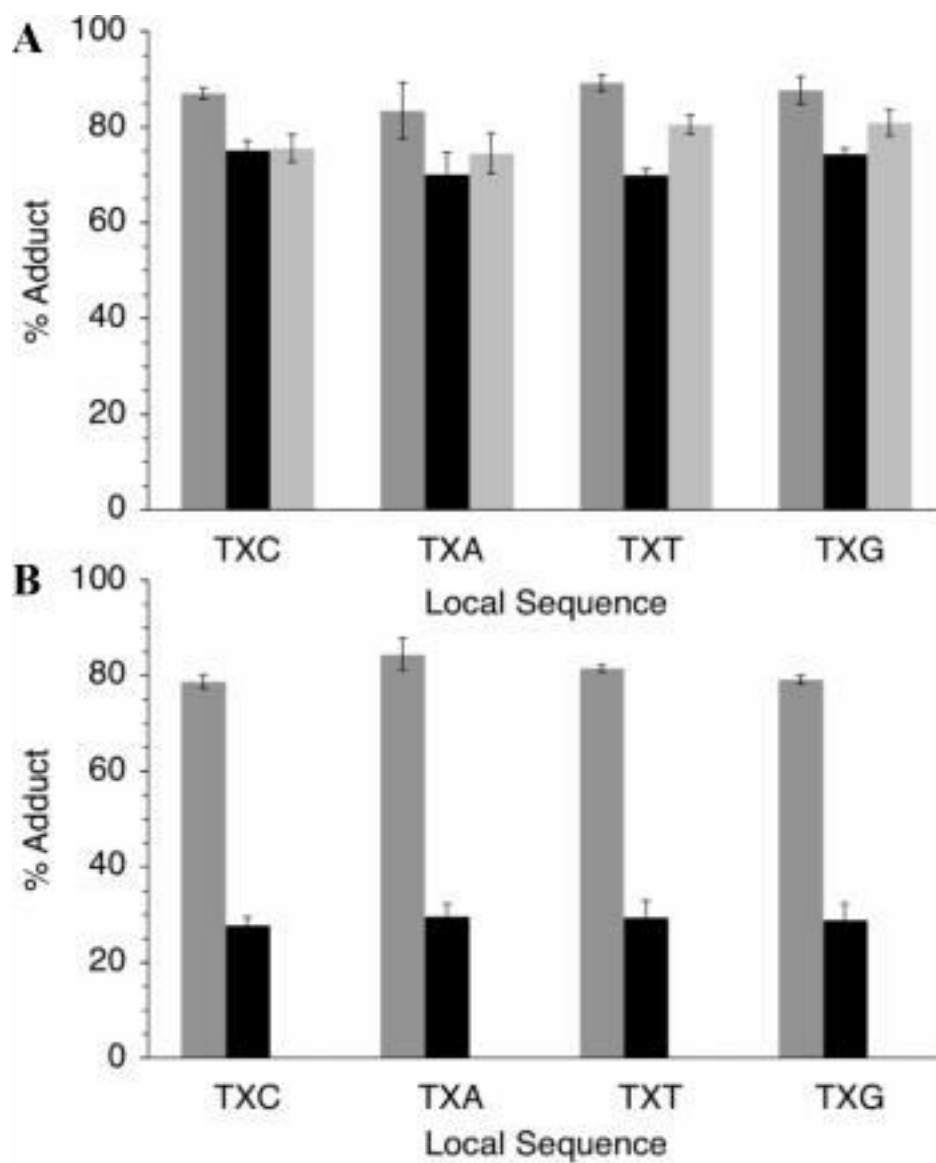
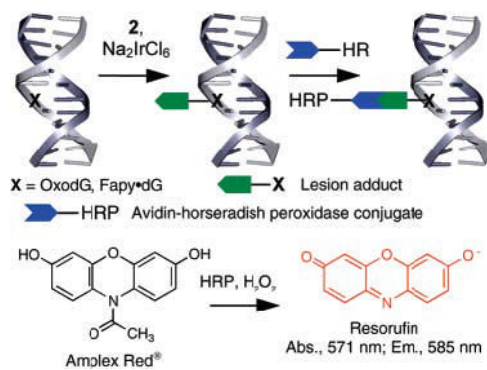


Figure 2. Average adduct yield with **2** in DNA as a function of sequence. (A) OxodG (**3a-d**). (B) Fapy•dG (**4a-d**). Dark gray, Na₂IrCl₆; black, Na₂IrBr₆; light gray, K₃Fe(CN)₆.



Scheme 1.
Fluorescence Detection of OxodG and Fapy•dG

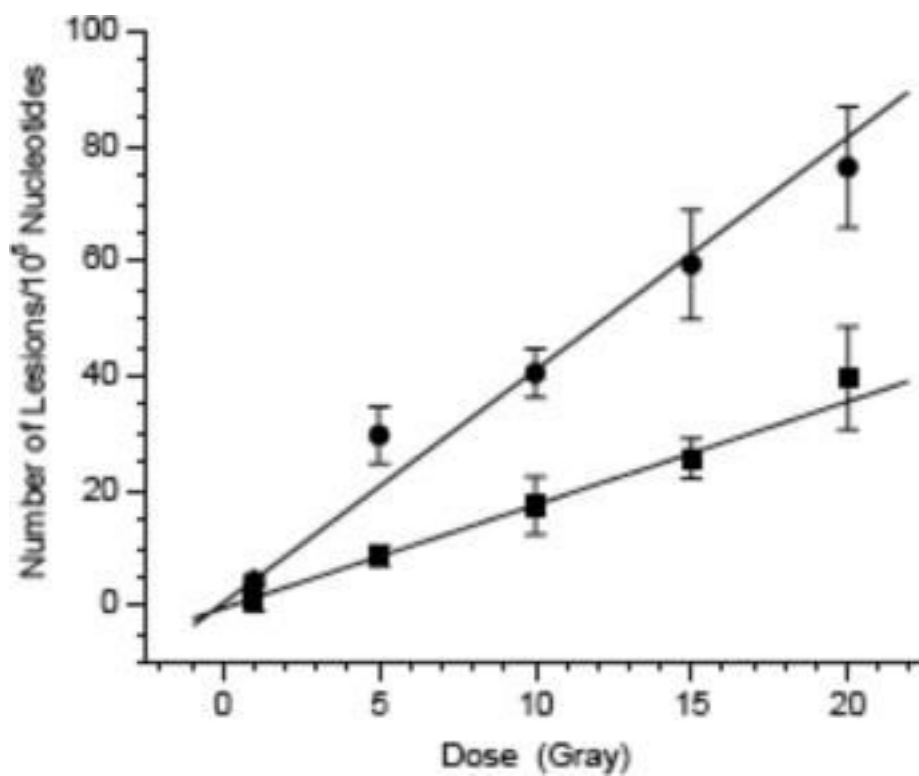


Figure 3. Average adduct yield with **2** in 287 nt duplex DNA (2.5 pmol) as a function of ^{137}Cs dose. Circles, Na_2IrCl_6 ; squares, $\text{K}_3\text{Fe}(\text{CN})_6$.