Effect of Low-Energy Electron Irradiation on DNA Damage by Cu^{2+} Ion

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ABSTRACT

Background: The combined effect of the low energy electron (LEE) irradiation and Cu^{2+} ion on DNA damage was investigated.

Materials and Methods: Lyophilized pBR322 plasmid DNA films with various concentrations (1-15 mM) of Cu^{2+} ion were independently irradiated by monochromatic LEEs with 5 eV. The types of DNA damage, single strand break (SSB) and double strand break (DSB), were separated and quantified by gel electrophoresis.

Results and Discussion: Without electron irradiation, DNA damage was slightly increased with increasing Cu ion concentration via Fenton reaction. LEE-induced DNA damage, with no Cu ion, was only 6.6% via dissociative electron attachment (DEA) process. However, DNA damage was significantly increased through the combined effect of LEE-irradiation and Cu ion, except around 9 mM Cu ion. The possible pathways of DNA damage for each of these different cases were suggested.

Conclusion: The combined effect of LEE-irradiation and Cu ion is likely to cause increasing dissociation after elevated transient negative ion state, resulting in the enhanced DNA damage. For the decrease of DNA damage at around 9-mM Cu ion, it is assumed to be related to the structural stabilization due to DNA inter- and intra-crosslinks via Cu ion.

Keywords: DNA damage, Copper ion, Electron beam, Dissociative electron attachment

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Introduction

Since Sanche and the collaborators have pioneered the investigation of DNA damage induced by low-energy electrons (LEEs) irradiation [1], it has been established that LEEs can cause DNA damages such as single-strand break (SSB), double-strand break (DSB), base lesion, and cross-links between nucleobases. LEEs of having energies not enough to ionize DNA molecules can react with DNAs and their subunits and lead to various DNA damages indirectly. For example, Boudaiffa et al. [1] have shown that LEEs with less than 15 eV can induce DNA damages on DNA films and explained the observation with dissociative electron attachment (DEA) process [2-8]. However, DNA *in vivo* is surrounded by many other molecules such as proteins and metal ions as well as H_2O and interacts with these molecules. Among these interactants, metal ions are important catalysts for oxidative damaging reactions in DNA and other biomolecules [9-13]. For example, it is well established that oxidative DNA damage in the presence of Fe ions in the solution has occurred via Fenton reactions [14, 15]. Motivated by these important

and independent roles of LEEs and metal ions, we have previously reported the combined effect of LEE-irradiation and Fe^{3+} ion on DNA damage [16]. Briefly summarizing the experiment, lyophilized pBR322 plasmid DNA films with various concentrations (0-7 mM) of Fe^{3+} ion were independently irradiated by monochromatic LEEs with 3 or 5 eV energies. After the irradiation, DNA- Fe^{3+} films were recovered and analyzed by the agarose gel electrophoresis. In the case of *non-irradiated* DNA- Fe^{3+} films, there was a little DNA damage (< 10%) for Fe^{3+} ion up to 7 mM concentration. In the case of irradiated DNA films without Fe^{3+} ion, again there was only a little DNA damage (< 3%). However, in the case of *electron-irradiated* DNA- Fe^{3+} films, DNA damage was enhanced considerably. A single effect of either Fe^{3+} ion or LEE-irradiation did not cause a significant DNA damage, but when these two components were combined DNA damage was significantly increased compared to the *sum* of the damages caused both by Fe^{3+} ion and by LEE-irradiation independently. Stimulated by this result, we have done the similar investigation with copper ion Cu^{2+} , another transition metal ion, to see how they behave similarly or differently from Fe^{3+} .

Both Fe and Cu ions are transition metal ion and they share some common characters while show some differences. Transition metal ions such as cobalt(II), copper(II) have different binding characteristics toward DNA. They bind almost exclusively by coordinating to the N7 position of purine, especially of guanine [17]. Copper has long been biocidal tools from the ancient era and is one of a relatively small group of metallic elements that are essential to human health [18]. Catalytic copper, because of its mobilization and redox activity, is believed to play a central role in the formation of reactive oxygen species that bind very fast to DNA, and produce damage by breaking the DNA strands or modifying the bases and/or deoxyribose leading to carcinogenesis [19].

Following the similar methods as done in the Fe^{3+} case, in this report we investigated the effects of LEE-irradiation and Cu^{2+} ion on DNA damage. The multilayered DNA- Cu^{2+} films were prepared by lyophilization technique and irradiated by 5 eV electrons. After LEE-irradiations, the irradiated DNA- Cu^{2+} films were analyzed by electrophoresis.

Materials and Methods

Plasmid DNA pBR322 was purchased from New England BioLabs Inc. This DNA molecule is a commonly used, double-stranded and supercoiled DNA with 4,361 bp in length

and 2.83×10^6 daltons in weight. Cupric chloride (CuCl_2) was purchased from Sigma-Aldrich Chemical Co. (St. Louis, MO). All other reagents and solvents were purchased from Sigma-Aldrich Chemical Co. and used as received without further purification. SYBR GREEN I ($10,000\times$) as a DNA dye was purchased from Invitrogen and the concentration of SYBR GREEN I was diluted to $100\times$ with dimethyl sulfoxide (DMSO) stock solution. Tantalum (Ta) plate was purchased from Goodfellow Cambridge Limited (Huntingdon, UK) and was cut into 12 mm $\times$ 12 mm square pieces for use as DNA film substrate.

To prepare dried DNA films on Ta plates, a home-made lyophilizer was used [16]. In treating plasmid DNA and CuCl_2 , nanopure water (distilled and deionized water ($18.2 \text{ M}\Omega \cdot \text{cm}$, 25°C ; ddH₂O)) was used. pBR322 DNA solution was prepared mixed with TE (Tris-HCl 10 mM, EDTA 1 mM, pH 8.0) buffer to dilute to a concentration of $0.05 \mu\text{g } \mu\text{L}^{-1}$, and CuCl_2 solutions mixed with H₂O to make eight different concentrations from 0 mM to 15 mM. The prepared pBR322 DNA solution and CuCl_2 solutions were individually mixed in a 0.5 mL microcentrifuge tube to make DNA- Cu^{2+} complex. Each DNA- Cu^{2+} mixture solution (with a volume of 9 μL) was carefully dropped on a chemically clean Ta plate. The dropped DNA- Cu^{2+} solutions were spread on the Ta plates to a circular shape with a 5 mm diameter. Immediately after, the solutions became frozen using liquid nitrogen and then transferred to the small vacuum chamber of the lyophilizing system. Water molecules in the DNA- Cu^{2+} solutions were extracted by the cryogenic sorption pump, and finally dried DNA- Cu^{2+} films with about 5 monolayers (ML) on the Ta plates were obtained. For each run of the experiments, eight DNA- Cu^{2+} films were prepared and loaded into another vacuum chamber for LEE-irradiation. To make the DNA films without Cu ion, the similar procedure was followed.

The figures and the details of an experimental setup for LEE-irradiation on DNA films were given in the previous report [16]. In brief, it consists of an electron gun (E-gun), a Faraday cup, and a sample holder. The E-gun can generate a monoenergetic electron beam having 0.5 eV resolution over the entire sample surface. Each DNA film was individually irradiated with constant irradiation time, beam current, and incident electron energy. For 5-minute irradiation time, it was estimated a total exposure of approximately 1.8×10^{14} electrons per DNA film. The irradiation of DNA film was carried out under ultra-high vacuum of about 10^{-9} torr.

Gel electrophoresis was used to identify and quantify DNA

damage induced by LEE-irradiation and Cu^{2+} ion. The DNA films were recovered from Ta plates with 9 μL of TE buffer. After that, 1 μL of SYBR GREEN 1 (100 $\times$) and 2 μL of loading buffer (6 $\times$) were added to the recovered DNA solution. Thus, prepared DNA samples were loaded with DNA marker molecule onto 1% agarose gel submerged under TAE solution (1 $\times$). After electrophoresis at 50 V for 35 minutes, separated DNA bands were photographed using gel-documentation system (ImageQuent300, GE Healthcare, Buckinghamshire, UK), and each type of DNA damage was quantified using ImageQuent program. All data shown in the figures and the table indicate an average values with corresponding standard deviations of four to eight independent experiments.

Results and Discussion

The structure of supercoiled DNA can be changed to circular or linear DNA as the results of strand breaks, which allows us to identify the types of DNA damage induced by LEE-irradiation and Cu^{2+} ion using gel electrophoresis. The initial state of plasmid pBR322 DNA is mostly supercoiled-type with less than 5% nicks. Figure 1A shows the results for the gel electrophoresis of DNA damages induced by LEE-irradiation with 5 eV on the DNA- Cu^{2+} films as a function of Cu^{2+} concentration (1-15 mM). The solid squares indicate the loss of supercoiled DNA induced by Cu^{2+} ion concentrations without 5 eV electron irradiation and the solid circles indicate the loss of supercoiled DNA for the combined effect of 5

Fig. 1. (A) Loss of supercoiled DNA by the combined effect of LEE-irradiation and Cu^{2+} ion. The solid circles indicate the results for 5 eV electron irradiations on DNA- Cu^{2+} complex and the solid squares indicate the result without electron irradiation, respectively. Error bars at 0 mM are unseen because they are smaller than the data points. (B) Loss of supercoiled DNA by the combined effect of LEE-irradiation and Fe^{3+} ion. The solid circles indicate the results for 5 eV electron irradiations on DNA- Fe^{3+} complex and the solid squares indicate the result without electron irradiation, respectively [16].

Table 1. Yields of Total Damage for Non-irradiated and 5 eV Electron-irradiated DNA- Cu^{2+} Complexes

Cu^{2+} concentration (mM)	SC loss due to Cu^{2+} only [No e-irradiation]	SC loss due to Cu^{2+} and 5 eV electron	Enhancement due to combined effect of Cu^{2+} and 5 eV electron
0 [No Cu^{2+}]	5.61 ± 0.59	12.16 ± 0.90	6.55 [a]
1	4.88 ± 1.29	12.18 ± 6.33	0.75
3	6.25 ± 0.65	21.73 ± 5.92	8.93
5	10.20 ± 1.78	27.45 ± 9.28	10.70
7	9.02 ± 4.07	25.88 ± 12.14	10.31
9	7.77 ± 3.54	15.31 ± 4.18	0.98
11	11.24 ± 4.32	29.32 ± 16.89	11.53
13	13.35 ± 1.93	53.89 ± 15.89	33.99
15	24.34 ± 3.82	73.74 ± 17.89	42.85

All values in the last column, except value [a] = (Loss of SC DNA with LEE and Cu^{2+}) – (Loss of SC DNA with LEE only) – (Loss of SC DNA with Cu^{2+} only). Value [a] = (Loss of SC DNA with 5 eV irradiation) – (Loss of SC DNA with no LEE irradiation).

Fig. 2. The schematic drawings of intra- and inter-crosslinks between Cu ion and guanine bases. G-Cu-G intra-crosslink (A) and CG-Cu-GC inter-crosslink (B) show 2-D and 3-D images of G-G intra- and inter-crosslinks mediated by Cu ion, respectively.

eV electron irradiation and Cu^{2+} ion concentrations, respectively. Again, each point is an average of four to eight independent measurements with an error bar indicating the corresponding one standard deviation. In Table 1, the same results are tabulated for damage yields due to Cu^{2+} concentrations only and the combined effect of LEEs and Cu^{2+} ion. Each yield value is, as in the figure, expressed as an average percentage loss of supercoiled DNA, given with a corresponding standard deviation.

For the convenience of the comparison, the result for Fe^{3+} ion is reproduced in Figure 1B. In our original work on Fe ion, there are results for the electron energies of 3 eV and 5 eV [16]. However, in Figure 1B, only the result for 5 eV is given to compare with the Cu ion data. The general trend observed in Cu ion data looks similar to that of Fe ion, except in Cu ion case there is a prominent dip at 9 mM concentration for the irradiated data. As can be seen in Figure 1A and Table 1, when only Cu^{2+} ion was added to supercoiled DNA (i.e., without LEE-irradiation), there was only a slight damage for the Cu^{2+} concentration range of 0 to 9 mM and slowly started to increase from 11 mM (see the solid square in Figure 1A). This damage might be generated by Fenton reaction of Cu^{2+} ion and H_2O_2 during the film preparation as previous Fe^{3+} case [20, 21]. In the case when there was only LEE-irradiation

on DNA with no Cu^{2+} ion, a little DNA damage of about 6.6% has occurred. In other words, a single effect of either Cu^{2+} ion or LEE-irradiation did not cause a significant damage. However, when these two components were combined, the DNA damage was increased quite significantly with increasing Cu^{2+} ion concentration except around 9 mM, compared to the sum of DNA damages caused by Cu^{2+} ion and by LEE-irradiation independently. At 5 eV electron energy, most of DNA damage has been identified as SSB in the spectra of gel electrophoresis. In the previous experiment with Fe ion, we have done the experiments for the Fe^{3+} concentration from 0 to 7 mM, and above 7 mM there was a sharp increase of damage which might be due to an extra mechanism other than Fenton reaction [16]. But in copper case, the beginning of damage decrease for the electron-irradiated samples was noticed, and therefore the measurements with Cu^{2+} ion concentration up to 15 mM were made to observe and locate the dip around 9 mM.

As mentioned before, the overall trend of DNA damage by the combined effect of LEEs and Cu ion is similar to that of Fe ion. However, an interesting new observation in this study is a decreased DNA damage by 5 eV electron irradiation at around 9 mM Cu^{2+} ion. It is believed that, at the Cu^{2+} ion concentration near 9 mM, the chemical structure of DNA- Cu^{2+}

Fig. 3. The reaction pathways of DNA damage generated by single and combined effects of LEE-irradiation and Cu^{2+} ion. Path (A) and (B) show the fundamental DEA process, the combined effect of DEA and Fenton reaction and structural stabilization, respectively.

ion complex become stable to protect DNA from the damage by LEEs. Copper was found to bind DNA with high affinity and a crystal structure was formed between CuCl_2 and DNA giving a copper-binding to N7 of guanine residue and forming a pseudo-octahedral geometry [22, 23]. It is pointed by Gao et al. [17] that Cu(II) ion can bind exclusively to N7 of guanines and is capable of pulling two guanines (G10 and #G12 of CGCGTG) closer by about $1.5 \text{ \AA}$ to form a bis-coordination linkage. This suggests that Cu(II) ion may be very effective in causing inter- or intra-crosslinks between two different DNA duplexes, at least in the case of Z-DNA. Hay and Morris [24] proposed that copper may stabilize the helix *via* a charge transfer complex formed when copper, acting as an electron acceptor, intercalates between two adjacent G-C pairs which act as electron donors. Indeed, copper has been shown to bind preferentially to G-C pairs. This kind of stabilization may occur at a certain Cu^{2+} concentration and protect DNA from being damaged by LEEs. From the basis of these points, we suggest the possible chemical binding structures of guanines and Cu^{2+} ion. In Figure 2A, the neighboring guanines in a single strand can be crosslinked via making bonding with Cu ion. This intra-crosslink leads to a bending structure between guanine bases. In Figure 2B, the close opposite guanines in a double strand can also be crosslinked via making bonding with Cu ion. This inter-crosslink leads to a pulling structure between the guanine bases on

the complementary DNA strands. In other words, the chemical structure of DNA duplex becomes stable via bending and pulling effects of intra- and inter-crosslinks on the single and double strands, respectively. This structural change might be capable of reducing the DEA processes due to the change of resonance energy level for capturing LEEs. Consequently, this leads to the decrease of DNA damage near 9 mM Cu ion.

The reaction pathways of DNA damage generated by single and combined effects of LEEs and Cu^{2+} ion are proposed in Figure 3. The fundamental damage process is DEA via LEE-irradiation as seen path (A) in Figure 3. This process has been well known and generally been accepted to explain DNA damage by collisions of LEEs (below 10 eV) [1, 16]. In the presence of Cu^{2+} ion, guanine base in DNA strand can preferentially interact with Cu^{2+} ion to form DNA- Cu^{2+} complex. Fenton reaction is main cause of DNA damage without LEE-irradiation, and this damage is generally less than that induced by LEEs. LEE-irradiation on DNA- Cu^{2+} complex is capable of increasing DNA damage through the elevated DEA process. However, in a certain concentration of Cu^{2+} ion (9 mM in this experiment), DNA damage by LEE-irradiation is conversely decreased due to structural stabilization. These processes are summarized in Figure 3 path (B). To fully understand DNA damage or protection mediated by LEEs and transition metal ions, it requires more experimental and theoretical evidences.

Conclusion

The low energy electron-induced DNA damage in the presence of Cu^{2+} ion was investigated. The single effect of LEE collision or Cu^{2+} ion on the yield of DNA damage does not exceed 20%, while the combined effect of two causes was significantly increased up to 43% as a function of Cu^{2+} ion concentration. Interestingly, a protection-like effect at around 9 mM Cu^{2+} ion on LEE-induced DNA damage was observed. The possible DNA damage pathways for the mechanism involved was suggested. For fully understanding whole mechanisms, it needs further studies.

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