Preventive Effects of Active Hexose Correlated Compound (AHCC) on Oxidative Stress Induced by Ferric Nitrilotriacetate in the Rat

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SUMMARY
Ferric nitrilotriacetate (Fe - NTA) is a strong oxidant, which generates highly reactive hydroxyl radical and causes injuries of various organs including the kidney and liver. The formation of 8- hydroxy - 2' - deoxyguanosine (8- OHdG) adducts in the renal DNA is one of the earliest events after treatment with Fe - NTA. Since Active Hexose Correlated Compound (AHCC), an extract of fungi, has been shown to act as an antioxidant, its protective effect on the oxidative stress induced by Fe - NTA was examined in the present study. AHCC at 3% in drinking water was given to male Wistar rats for 1 week, then Fe - NTA was injected intraperitoneally. At 3 h after the treatment with Fe - NTA, levels of 8- OHdG in the bladder urine, creatinine in the serum, thymic apoptosis, serum levels of aspartate and alanine aminotransferases were significantly increased. All of these increases were restored to normal by the AHCC pretreatment. These results suggest that AHCC is potent in restoring the disorders of various organs induced by oxidative stressors.

Key Words: ferric nitrilotriacetate, 8- hydroxy - 2' - deoxyguanosine, creatinine, thymic apoptosis, aminotransferases, AHCC

INTRODUCTION
Reactive oxygen species (ROS) including superoxide (O2−), hydrogen peroxide (H2O2), hydroxyl radical (·OH) are constantly generated in aerobic respiration in response to both external and internal stimuli. Low levels of ROS play an important role in signal transduction, cell proliferation, apoptosis, immunity and defense mechanisms. High levels of ROS, however, cause severe metabolic malfunctions and damage to biological macromolecules. For example, excess ROS induce oxidative damages to nucleic acids and proteins, decrease the efficiency of DNA polymerase repair and activity of signal transduction and cause cell death by apoptosis or necrosis. ROS induce chemical changes in bases and change in DNA conformation, causing mutation. Damage to DNA by ROS is critical for carcinogenesis. Moreover, the accumulation of free radical - mediated damage in DNA is the major cause of the physiological changes associated with aging.

Animals including human possess numerous machineries of antioxidative defense by themselves. Such defense mechanisms within the organism have evolved to limit the levels of reactive oxidants and the damage that they inflict. Enzymes involved in such mechanisms include superoxide dismutase (SOD), glutathione peroxidase and catalase. Moreover, there exist many structural defenses, for example, sequestration of H2O2 - generating enzymes in peroxisomes and chelation of any.

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free iron or copper salts in transferrin and ferritin or ceruloplasmin to avoid the product of Fenton reaction, 
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Epidemiological studies have suggested that antioxidative supplements, such as dietary antioxidants of small molecules, including vitamin C (ascorbate), vitamin E (tocopherol) and carotenoids are potent in preventing carcinogenesis 10, 11. Thus, natural plant components have been expected to provide oxidative defenses against cancer and degenerative diseases caused by oxidative stress. Active Hexose Correlated Compound (AHCC™ Amino UP Chemical Co. Ltd. Sapporo) is a mixture of polysaccharides, amino acids and minerals derived from fungi. It is obtained by hot water extraction after culturing mycelia of several basidiomycetes in a liquid culture tank and treating them with several enzymes 12. The main active component of AHCC is a mixture of oligosaccharides whose average molecular weight is approximately 5,000 12, 13. The chemical analysis has revealed that polysaccharides are the major components of AHCC, consisting approximately 74%. Nearly 20% of the polysaccharides are α-1, 4-glucan and its acetylated forms, which are thought to be active components 12. AHCC also contains some other polysaccharides including β-glucans. Both a- and β-glucans are shown to be responsible for the antitumor effects of Basidiomycetes 14. It has been reported that this extract shows various beneficial effects in humans and in experimental animals acting either as a biological response modifier (BRM) or an antioxidant. It was shown that AHCC was effective in suppressing growth of experiment cancers in mice and rats such as SST-2 tumor and breast cancer 15. AHCC enhances a helper T cell 1 (Th1) response in susceptible BALB/c mice, especially increasing interleukin (IL) - 12 levels 15. It was also found out that Th1 cytokines levels such as tumor necrosis factor (TNF) - α and interferon (IFN) - γ were increased in response to AHCC treatment. AHCC is used clinically as a successful complementary therapy for cancer patients 16. AHCC treatment after surgical operation could improve the quality of life (QOL) of patients with malignant tumors and increase the survival rate by inhibiting tumor metastasis 16. AHCC can also prevent the onset of the diabetes induced by streptozotocin 17. Recently, we have found that AHCC suppresses thymic apoptosis induced by dexamethasone 18, suggesting that AHCC acts as an antioxidant.

Ferric nitrotriacetate (Fe - NTA) is a chemical that induces severe oxidative damage by Fenton reaction, in which the very reactive hydroxyl radical is generated19-23. It acts as a potent nephrotoxin, causing high incidence of renal cell carcinoma in rats and mice 6, 20, 21. Fe - NTA treatment results in renal DNA damage, including formation of 8 - hydroxy - 2' - deoxyguanosine (8-OHdG) in the renal DNA through the generation of active oxygen radicals. The increase in urinary excretion of 8-OHdG reflects the oxidative DNA damage in vivo 19, 23. Creatinine levels in the serum are significantly increased following Fe - NTA treatment 19, 23i. Fe- NTA also can induce hepatic injuries acting through the generation of ROS 20. These damages done to several organs are ameliorated by antioxidants 27, 28.

The objective of our present study is to examine whether or not AHCC has the preventive effects on the damage induced by oxidative stress in rats treated with Fe - NTA. Our present study deals with the beneficial effects of AHCC on the damage of various organs induced by Fe - NTA.

**MATERIALS AND METHODS**

1. Materials. Lyophilized AHCC was prepared by Amino UP Chemical Co. Ltd. (Sapporo, Japan). Fe -NTA solution was prepared immediately before use as previously described22. Nuclease P1 was obtained from Sigma (St. Louis, MO, USA).

2. Animals. Studies with the rats were approved by Animal Care and Use Committee, Dokkyo University School of Medicine. The animals were treated according to guidelines for the Care and Use of Laboratory Animals of the Committee. Male Wistar rats of 8 - weeks old were purchased from Charles River Japan Inc. (Kanagawa, Japan). The animals were kept in a room at 23 ± 2°C with 12 h light and dark cycle and kept free access to food and water.

Twenty- one rats were divided into four groups. Two groups received 3% AHCC as a drinking water for a week. The concentration of AHCC and the duration of the treatment were chosen according to our previous results 18. Control groups received only tap water for a week. Fe - NTA (15 mg/kg body weight as Fe$^{3+}$ was injected intraperitoneally to two groups; one of them
was AHCC - pretreatment group, and the other was the control group. Fe-NTA effects at this dose were maximal at 3 h of the treatment as shown by others. 3. Blood sampling and organ preparation. Rats were sacrificed by decapitation at 3 h after Fe-NTA-treatment. Blood samples were collected into sampling tubes. Serum samples were obtained after centrifuging at 4°C for 15 min at 3,000 rpm. The sera thus obtained were used for the assay of serum aminotransferases and creatinine.

Thymus glands were removed and washed in cold phosphate-buffer solution (PBS), then put into the tubes which contained collagenase-EDTA for apoptosis assay. Urine samples were taken from urinary bladder of rats for 8-OHdG assay.

4. 8-OHdG assay in the urine. Urine samples were centrifuged at 10,000 rpm, and the supernatant was used for ELISA analysis. 8-OHdG was assayed by ELISA kits, which were prepared by Japan Institute for the Control of Aging (Shizuoka, Japan).

5. Detection of apoptosis by flow cytometry. Cell suspensions were obtained after collagenase-EDTA (0.25% in PBS) treatment and filtration through nylon mesh. The cell suspensions were washed with PBS (pH 7.4), adjusted to a concentration of 1.5 x 10^6 cell/mL. Then the cells were centrifuged at 200 x g for 5 min. The 200 x g centrifuged cell pellet was gently resuspended in 1.5 mL hypotonic fluorochrome solution containing propidium iodide (PI, 50 μg/mL in 0.1% sodium citrate plus 0.1% Triton X 100), in polypropylene tubes. The tubes were placed at 4°C in the dark overnight before the flow cytometric analysis. Flow cytometry was carried out by analyzing 10,000 cells per test using a FACSCalibur (Becton Dickinson, Mountain View, CA, USA).

6. Creatinine and aminotransferase assays. Creatinine in the serum was assayed by creatinine kit. The activity of alanine aminotransferase (ALT), aspartate aminotransferase (AST) in the serum were routinely assayed by commercial kits.

7. Statistical analysis. All the data were expressed as mean ± SEM. Analysis of variance (ANOVA) was performed and Scheffe's multiple comparison test was applied to test for the differences between individual groups. A p value less than 0.05 was considered statistically significant.

RESULTS

1. 8-OHdG in the bladder urine

First we observed the noxious effects of Fe-NTA in the kidney and the effects of simultaneous administration of Fe-NTA and AHCC on the renal function. 8-OHdG in the bladder urine increased significantly at 3 h after treatment with Fe-NTA. A significantly lower level of 8-OHdG in the urine of AHCC pretreated rats was observed, when compared with that of the rats treated with Fe-NTA alone (Fig. 1).

2. Creatinine in the serum

The enhanced serum creatinine levels, which depend on glomerular filtration rate, are indicative of renal injury22, 23. In our experiment, the creatinine levels in the serum increased significantly in the rats treated with Fe-NTA treatment alone, while AHCC - pretreatment restored the increased levels to normal (Fig. 2).

4. Serum AST and ALT

Hepatic injuries induced by Fe-NTA were assessed by measuring the activity of serum aminotransferases. The serum levels of AST and ALT were elevated in the Fe-NTA-treated group than those in the control and AHCC groups (Figs. 3, 4). AHCC - pretreatment decreased AST and ALT levels in the serum significantly to the normal levels.

3. Thymic apoptosis

Thymic apoptosis in the Fe-NTA treated rats increased significantly compared with that in the control and AHCC groups. Thymic apoptosis in the Fe-NTA - AHCC treated rats was much lower than in the rats treated with Fe-NTA alone (Fig. 5).

DISCUSSION

The results of the present study have shown that Fe-NTA increases the urinary levels of 8-OHdG, and serum levels of creatinine, AST and ALT. All these changes induced by Fe-NTA have already been observed by others, showing that this toxin damages both the kidney and the liver16, 22, 23. In the present study it was first shown that thymic apoptosis was induced by this chemical. These changes may be closely associated with oxidative stress, because Fe-NTA produces ROS in vivo. All the Fe-NTA - induced changes observed here are restored to normal by the pretreatment with AHCC for 1 week. The results seem to suggest that AHCC can protect
Fig. 1: Effects of Fe-NTA and AHCC on 8-OHdG levels in urinary bladder urine. Urine samples in the urinary bladder were withdrawn immediately after sacrifice. Fe-NTA (15 mg Fe $^{3+}$/Kg) was injected i.p. 3h before experiment. AHCC at 3% in drinking water was given for a week. Data are shown as means ± SEM with the numbers of determination in parentheses.

*, p< 0.001 vs. control group.
a, p< 0.001 vs. AHCC + Fe-NTA group.
Abbreviations are as follows: FNT, Fe-NTA, ferric nitrilotriacetate; AHCC, Active Hexose Correlated Compound and 8-OHdG, 8-hydroxyl-2'-deoxyguanosine.

Fig. 2: Effects of Fe-NTA and AHCC on serum creatinine levels. Creatinine in the serum were assayed by creatinine kit. Data are shown as means ± SEM with the numbers of determination in parentheses.

*, p< 0.001 vs. control group.
a, p< 0.001 vs. AHCC + Fe-NTA group.
Fig. 3: Effects of Fe-NTA and AHCC on serum alanine aminotransferase (ALT) levels. ALT was assayed by commercial kit. Data are shown as means ± SEM with the numbers of determination in parentheses.

*, p < 0.001 vs. control group.

a, p < 0.001 vs. AHCC + Fe-NTA group.

Fig. 4: Effects of Fe-NTA and AHCC on serum aspartate aminotransferase (AST) levels. AST was assayed by commercial kit. Data are shown as means ± SEM with the numbers of determination in parentheses.

*, p < 0.001 vs. control group.

a, p < 0.001 vs. AHCC + Fe-NTA group.
Fig. 5: Effects of Fe -NTA and AHCC on the thymic apoptosis.

Apoptosis was estimated by flow cytometry using FACSCalibur. Data are shown as means ± SEM with the numbers of determination in parentheses.

- *p < 0.001 vs. control group.
- a, p < 0.001 vs. AHCC + Fe-NTA group.

Fe-NTA is a renal toxicant and carcinogen in rats and mice. This chemical increases markedly the urinary excretion of 8-OHdG which is a useful marker for measuring the level of oxidative DNA damage. The damage in DNA is closely associated with aging and carcinogenesis.

A common form of DNA damage is the formation of hydroxylated bases, which is considered to be an important event in carcinogenesis induced by oxidative stress. The formation of 8-OHdG has been shown to be suppressed by antioxidants such as vitamin E, vitamin C and methionine. Many other antioxidants have been shown to protect the injuries induced by Fe -NTA. For example, N-acetylcysteine (NAC), a precursor of intracellular cysteine and glutathione, prevents renal damage induced by Fe -NTA in the rat.

Alpha- tocopherol, a lipid-soluble antioxidant, ameliorates renal proliferative response and toxicity induced by Fe -NTA. AHCC decreased the Fe -NTA - increased 8-OHdG levels in the urine, suggesting that this compound prevents mutation by scavenging ROS.

It is reported that Fe -NTA increases the hepatic ornithine decarboxylase (ODC) and that the pretreatment of rats with butylhydroxytoluene (BHT), an antioxidant, suppresses the increase in the enzyme activity. Fe -NTA injected intraperitoneally into rats increases plasma levels of the two aminotransferases, ALT and AST and AHCC pretreatment normalized the elevated enzyme activities. These findings suggest that AHCC protect the hepatic damage induced by oxidative stress of Fe -NTA.

Thymic apoptosis is induced by glucocorticoids and oxidative stress and suppressed by antioxidants such as melatonin. Recently we have found that AHCC suppresses thymic apoptosis induced by dexamethasone. All these findings suggest that AHCC acts as an antioxidant. It is not known yet if AHCC acts directly as an antioxidant to scavenge ROS or induces some enzymes, which cleavage ROS. It still remains unclarified which components of AHCC are responsible for its protective effect against oxidative stress. Our preliminary experiments have revealed that AHCC itself has no antioxidant effect on thymocytes in vitro (data not shown). The study is currently under way to test the effects of individual α- and β-glucans purified from AHCC.

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