Protective Effects of Vitamin E Analogs against Carbon Tetrachloride-Induced Fatty Liver in Rats

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Summary Recently, it has been reported that α-tocopherol (α-Toc) is effective for amelioration of liver damage. However, it is unknown whether other vitamin E analogs are effective. In this study, we investigated the effects of γ-tocopherol (γ-Toc) and tocotrienols (T3) in rats with fatty liver. Rats fed a vitamin E-deficient diet for four weeks were divided into eight groups: Control, carbon tetrachloride (CCl₄), α-Toc, α-Toc + CCl₄, γ-Toc, γ-Toc + CCl₄, T3 mix, T3 mix + CCl₄. After a 24 h fast, the rats were administered 20 mg of each of the vitamin E analogs, respectively. Moreover, the CCl₄ group were given 0.5 ml/kg body weight corn oil preparation containing CCl₄ 6 h after vitamin E administration. We measured the activities of aspartate aminotransferase and alanine aminotransferase (ALT) in plasma, and the contents of triglyceride (TG), total cholesterol (T-Chol) and vitamin E analogs in the liver. Also, we determined the hepatic expression of mRNA for inflammatory cytokines. The liver TG content in the γ-Toc + CCl₄ and T3 mix + CCl₄ groups was decreased in comparison with the CCl₄ group. Moreover, ALT activity in the T3 mix + CCl₄ group was significantly lower than CCl₄ group. These findings suggest that γ-Toc and T3 are effective for amelioration of fatty liver.

Key Words: γ-tocopherol, tocotrienol, fatty liver, carbon tetrachloride

Introduction

Carbon tetrachloride (CCl₄) is a chemical agent widely used for experimental induction of fatty liver and liver fibrosis in animals [1]. It is considered that CCl₄ is metabolized by cytochrome P-450 (CYP) to unstable trichloromethyl free radicals (e.g. CCl₃, CCl₃O₂⁻), which then bind covalently to membrane proteins, finally causing lipid peroxidation [1–4]. The activities of aspartate aminotransferase (AST) and alanine aminotransferase (ALT) in rat plasma rise remarkably with hepatocyte necrosis and lipid accumulation when CCl₄ is administered to rats. Therefore, this model is used widely to induce various fatty liver states.

Vitamin E is present naturally in the form of eight analogs, each differing in the number and position of the methyl groups on the chroman ring and by the presence or absence of a double bond on the phytol side chain. One group, the tocopherols (Tocs), have a saturated phytol side chain, and the other, the tocotrienols (T3s), have an unsaturated phytol side chain. Each of the groups can occur in α-, β-, γ-, or δ-forms. Especially, α- and γ-Toc are abundant in vegetable oil and seeds such as almonds and peanuts, and act as lipid-soluble antioxidants in cell membranes, being important for maintenance of cell membrane fluidity. When administered orally, Toc analogs (especially α- and γ-Toc) are equally absorbed from the small intestine without discrimination. After uptake into intestinal cells, Toc analogs are integrated into chylomicrons. Chylomicron remnants are catabolized during circulation by lipoprotein lipase. After uptake of chylomicron remnants by the liver, Toc analogs are discriminated by α-Toc transfer protein (α-TTP) in the
liver, which transports α-Toc preferentially over γ-Toc into very-low-density lipoprotein (VLDL) during assembly [5]. Therefore, non-α-Toc (γ-Toc, T3) remains in the liver. Accordingly, we hypothesized that non-α-Toc may show function more effectively than α-Toc in the liver. Some investigators have reported the effect of α-Toc on liver injury induced by CCl₄. It has already been reported that administration of α-Toc or an α-Toc-enriched diet suppresses the rise of plasma ALT activity and concentration of bilirubin induced by CCl₄ [6, 7], whereas non-α-Toc does not. In the present study we investigated the preventive effects of γ-Toc and T3 mix against lipid peroxidation, and also AST activities in CCl₄-treated rats.

Experimental Procedures

Animals and diets

Male SD-IGS rats (six weeks old) were purchased from Charles River Japan, Inc. (Kanagawa, Japan). Forty rats were divided into eight groups: Control (C), CCl₄ (CCl₄), α-Toc (αT), α-Toc + CCl₄ (αT + CCl₄), γ-Toc (γT), γ-Toc + CCl₄ (γT + CCl₄), T₃ mix (T₃), T₃ mix + CCl₄ (T₃ + CCl₄). They were housed individually in stainless steel wire netting cages, and fed a commercial chow (Nippon Clea Co., Tokyo, Japan). All experiments were conducted in accordance with the guidelines for the care and use of laboratory animals at Kanagawa Institute of Technology.

Administration of vitamin E analogs and induction of liver damage by CCl₄

After 24 h of fasting, 20 mg of each vitamin E analog was administered to the rats. The purity of T₃ mix (α-T3; 37.8%, β-T3; 4.0%, γ-T3; 45.5%, δ-T3; 10.7%) was 98%. The purity of α- and γ-Toc were above 96.1% respectively. All vitamin E analogs were gifted by Eisai Food & Chemical Co., Ltd. (Tokyo, Japan). Six hours later, a mixture of CCl₄ and stripped corn oil (1:1, 1 ml/kg body weight; 0.5 ml/kg body weight as CCl₄) was administered orally to the rats. The control rats were given stripped corn oil alone as a placebo. After 6 h of CCl₄ administration, all the rats were killed under diethyl ether anesthesia, and arterial blood, liver, adipose tissue, kidney, and adrenal gland were removed for analysis.

The plasma was separated from blood cells by centrifugation at 3,000 rpm for 10 min. All tissues were immediately stored at −80°C until analysis for biological parameters. Liver samples to be used for RNA isolation were soaked in RNA later solution immediately and stored at −80°C. Liver samples to be used for histopathological examination were fixed in 10% formalin.

Quantitative analysis of vitamin E analogs using HPLC

The quantity of vitamin E in each organ was measured using Ueda’s method [8]. A 0.1 g sample of each organ was homogenized with 0.9 ml of 0.9% NaCl (weight/vol) solution. The resulting homogenate (0.1 ml) was pipetted into a 10 ml centrifuge tube, and 50 μl of 1,2,2,5,7,8-pentamethyl-6-hydroxychroman (PMC, 1 μg/ml) as an internal standard and 1.0 ml of ethanolic pyrogallol (6%, weight/vol) were added to each tube with stirring. After 0.2 ml of 60% (weight/vol) KOH solution had been added to each tube, the contents were saponified at 70°C for 30 min. After cooling, vitamin E analogs were extracted with 4.5 ml of 1% sodium chloride solution and 3.0 ml of 10% ethyl acetate/n-hexane solution, and centrifuged at 3,000 rpm at 4°C for 5 min. A 2.0 ml aliquot of the upper layer was evaporated, dissolved in 0.2 ml of n-hexane, and subjected to HPLC. The HPLC system consisted of a pump, degasser, column oven, and detector (LC-20AD, DGU-20A3, CTO-20A and RF-10AXL, Shimadzu Co., Kyoto, Japan). The analytical conditions were as follows: column, Capcell pak NH2 column (4.6 mm I.D. × 250 mm; Shiseido, Tokyo, Japan); mobile phase, n-hexan-isopropanol (98:2); flow rate, 1.0 ml/min; detection wavelength, 325 nm.

Measurement of liver damage markers and lipid concentration in plasma

AST and ALT activities and triglyceride (TG), total cholesterol (T-Chol) concentrations in plasma were measured using a biochemical autoanalyzer (CA-180; Furuno Electric Co., Ltd., Hyogo, Japan).

Measurement of liver lipid content

Total lipid in the liver was extracted by the method of Folch [9]. A 0.9 g sample of liver was ground with sodium sulfate and extracted with chloroform:methanol = 2:1, and then made up to 25 ml. A 125 μl aliquot of the extract was added to 125 μl of 1% TritonX-100 ethanol solution, followed by heating at 50°C overnight, and then dissolved in 15 μl of distilled water. TG and T-Chol concentrations in liver were measured using a biochemical autoanalyzer.

Measurement of mRNA expression of inflammatory cytokines in liver

Total RNA was extracted from liver tissues using RNeasy. The quantity and purity of the RNA were determined from the absorbance at 258/280 nm. Total RNA was reverse-transcribed into cDNA using a high-capacity RNA-to-cDNA kit in accordance with the manufacturer’s protocol. 7500 Fast Real-Time PCR system and real time PCR kit (TaqMan®

Vol. 47, No. 2, 2010
Gene Expression Assays, Applied Biosystems Japan Ltd., Tokyo, Japan) were employed based on the manufacture’s instruction. β-actin was used as an internal control. The content of the primer/probe mixture of inflammatory cytokines and β-actin is shown in Table 1, and PCR thermal cycling conditions are shown in Table 2.

### Statistical analysis

All data are expressed as the mean ± SD. Statistical analysis was performed by one-way ANOVA, followed by Bonferroni post hoc test. Statistical analyses were performed using Kaleida Graph ver. 4 for Windows (Hulinks Inc., Tokyo, Japan). Differences were considered to be significant at \( p<0.05 \).

### Results

#### Liver weight and lipid content

There were no significant differences in liver weight among the groups. The content of TG in liver was increased by administration of CCl\(_4\), but the increase was significantly suppressed by administration of T3. On the other hand, there were no significant differences in liver T-Chol content among the groups (Fig. 1). In addition, histopathological examination of the liver was performed after hematoxylin-eosin staining. Lipid accumulation in the liver was confirmed in the CCl\(_4\) group, αT + CCl\(_4\) group, γT + CCl\(_4\) group and T3 + CCl\(_4\) group after administration of CCl\(_4\), but that in the T3 + CCl\(_4\) group was lower than in the CCl\(_4\) group (Fig. 2).

#### Activities of liver damage markers and lipid concentrations in plasma

AST activity in plasma was increased by administration of CCl\(_4\), but this was unaffected by administration of vitamin E analogs. On the other hand, ALT activity also tended to increase after administration of CCl\(_4\). However, the rise in the plasma ALT level tended to be suppressed by γ-Toc administration, and T3 had a significant suppressive effect (Fig. 3). There were no significant differences in the TG and T-Chol concentrations in plasma among the groups (data not shown).

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**Table 1. Primer probe mixture of inflammatory cytokines and β-Actin**

| Assay ID       | Refseq   |
|----------------|----------|
| Rat TNF-α      | Rn00667869_m1 NM_031144.2 |
| Rat IL1-β      | Rn00667869_m1 NM_031144.2 |
| Rat IL6        | Rn00667869_m1 NM_031144.2 |
| Rat β-actin    | Rn00667869_m1 NM_031144.2 |

**Table 2. Thermal cycling condition**

| Step 1 | Step 2 |
|--------|--------|
| Hold   | 40 Cycle |
| 95°C   | 95°C   |
| 20 s   | 3 s    |
| 30 s   |        |

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Fig. 1. Effects of vitamin E analogs on lipid contents of rat liver after administration of CCl\(_4\); (A) Liver TG content. (B) Liver T-Chol content. Rats were fed a vitamin E-deficient diet for four weeks. After a 24 h fast, 20 mg of each vitamin E analog was administered. Six hours later, a mixture of CCl\(_4\) and stripped corn oil (1:1, 1 ml/kg body weight; 0.5 ml/kg body weight as CCl\(_4\)) was orally administered to the rats. The control rats were given stripped corn oil alone as a placebo. Six hours after CCl\(_4\) administration, the liver was removed and its TG and T-Chol contents were measured. C; Control group, CCl\(_4\); CCl\(_4\) dosage group, αT; α-Toc dosage group, α-T + CCl\(_4\); α-Toc + CCl\(_4\) dosage group, γT; γ-Toc dosage group, γ-T + CCl\(_4\); γ-Toc + CCl\(_4\) dosage group, T3; T3 mix dosage group, T3 + CCl\(_4\); T3 mix + CCl\(_4\) dosage group. The values are mean ± SD for 5 rats, *\( p<0.05 \), **\( p<0.001 \) (one-way ANOVA followed by Bonferroni post hoc test).
Expression of mRNA for inflammatory cytokines in liver

Expression of mRNA for tumor necrosis factor alpha (TNF-α) and interleukin-1 (IL1)-β in the liver was increased by administration of CCl₄ (Fig. 4), suggesting that CCl₄ administration caused inflammation of the liver parenchyma.

Vitamin E analog contents of various tissues

The expression of TNF-α and IL1-β mRNA was increased further when γ-Toc was administered in such a state.

The α-Toc content of liver and plasma was not changed.
by administration of CCl4. However, the α-Toc content of the liver showed a tendency to decrease. On the other hand, the α-T3 content of plasma was significantly decreased by CCl4 (Fig. 5). There were no significant differences in the quantity of vitamin E analogs in other tissues.

**Discussion**

In this study, we investigated the effects of γ-Toc and T3 on fatty liver induced by CCl4 in rats. At first, liver weight was increased in each group by administration of CCl4, thus confirming the induction of liver hypertrophy. As the TG content of the liver was also increased, this hypertrophy appeared to be caused by accumulation of TG. Because the increase in the liver TG content was suppressed by administration of T3, the latter may regulate lipid accumulation in the liver. Various hypotheses have been proposed to explain the mechanism of fatty liver induced by administration of CCl4. Recknagel et al. [10] suggested that CCl4 controls the secession of ribosome and liver proteosynthesis, and blocks lipoprotein secretion. In addition, the lipid accumulation is associated with deactivation of metabolizing enzymes. In fact, deactivation of various metabolic enzymes and a decrease of the P450 content in liver by administration of CCl4 have confirmed in both in vitro and in vivo systems [11, 12]. Because lipid accumulation in the liver was inhibited by T3 administration, T3 play a role in lipoprotein synthesis and the transportation of lipids, and may possibly inhibit the deactivation of metabolic enzymes.

It is known that AST and ALT activity in plasma is increased by acute liver damage. In particular, plasma ALT activity increases as a result of destruction and necrosis of hepatocytes. As ALT activity was inhibited by administration of T3, hepatocyte necrosis may have been suppressed. It has been reported that α-Toc ameliorates fatty liver damage induced by CCl4; ALT activity and the concentration of bilirubin were lower, and liver fibrosis was inhibited in rats fed an α-Toc-enriched diet [7]. In the present experiment, ALT activity tended to be lower in the γ-Toc group, and was significantly lower in the T3 group. Therefore, it is suggested that T3 suppresses the necrosis of hepatocytes by inhibition of CCl4-induced lipid peroxidation. It has reported that α-T3 possesses 40–60 times higher antioxidant activity against oxidative damage than α-Toc [13]. Consequently, we suggested that T3, especially α-T3, restrained strongly lipid peroxidation in the liver than α-Toc, and controlled lipid accumulation to the liver and cell death. A further examination is necessary for comparison of the effect of each T3 analog.

The α-T3 content of plasma was decreased by administration of CCl4, whereas the liver α-T3 content was unaffected. The α-Toc is preferentially transported by α-TTP in comparison with other vitamin E analogs, and is used for VLDL synthesis [5]. It is suggested that γ-Toc and T3 are accumulated in the liver and metabolized immediately. As it has been reported that CCl4 deactivates CYP, we considered that the metabolism of vitamin E may be interrupted by CCl4, and that T3 may remain in the liver for a long time. Consequently, we suggest that T3 may trap trichloro radicals in the liver, and thus show antioxidant activity.

Kupffer cells, which are fixed macrophages, are activated by liver damage, leading to production of inflammatory mediators.
cytokines such as TNF-α and IL1-β. This inflammation spreads by stimulation of the endothelium, and leads to localized migration of monocytes and neutrophils. IL-6 is also produced by T cells. In the present study, we found that the expression of mRNA for inflammatory cytokines was drastically increased by CCl₄. However, the vitamin E analogs T3, α-Toc and γ-Toc, did not affect the expression of inflammatory cytokine mRNAs. Because the expression mRNAs for inflammatory proteins changes markedly in states of acute liver damage, further examination of this issue will be necessary.

Recently, the prevalence of non-alcoholic fatty liver disease (NAFLD) has been increasing in Japan. Non-alcoholic steatohepatitis (NASH), one of the conditions of NAFLD, was first reported by Ludwig et al. [14]. It is considered that fatty liver progresses further to NASH under conditions such as oxidative stress and the action of inflammatory cytokines. The histopathological picture in patients with NASH resembles that of alcoholic steatohepatitis. However, this condition is harder to treat than alcoholic steatohepatitis, and can lead to liver cirrhosis or cancer. It has been reported that α-Toc is effective for NASH, and it has already been applied therapeutically. A report by Hasegawa et al. [15] indicated that ALT activity in the serum of NASH patients was reduced, and that the histopathological picture (fatty degeneration, inflammation, and fibrosis) was improved by administration of α-Toc (300 mg/day) for one year. Similar therapeutic application of γ-Toc and T3s is expected.

In summary, administration of vitamin E analogs has been shown to improve the liver damage induced by CCl₄ in rats, and it is suggested that the improvement effect of T3s is stronger than that of γ-Toc. From reports that T3s are effective for treatment of NASH, similar effects are expected for these other vitamin E analogs. A further experiment is necessary for comparison of the effect of each T3 analog. Moreover, detailed examinations employing other models are planned.

Fig. 5. Contents of individual vitamin E analogs in liver and plasma of rats administered CCl₄; (A) Liver α-Toc content. (B) Plasma α-Toc content. (C) Liver γ-Toc content. (D) Plasma α-T3 content. Rats were fed a vitamin E-deficient diet for four weeks. After a 24 h fast, 20 mg of each vitamin E analog was administered to the rats. Six hours later, a mixture of CCl₄ and stripped corn oil (1:1, 1 ml/kg body weight; 0.5 ml/kg body weight as CCl₄) was administered orally to the rats. The control rats were given stripped corn oil alone as a placebo. Six hours after CCl₄ administration, the liver and plasma were taken, and the quantity of each vitamin E analog was measured by HPLC. C; Control group, CCl₄; CCl₄ dosage group, αT; α-Toc dosage group, αT + CCl₄; α-Toc + CCl₄ dosage group, γT; γ-Toc dosage group, γT + CCl₄; γ-Toc + CCl₄ dosage group, T3; T3 mix dosage group, T3 + CCl₄; T3 mix + CCl₄ dosage group. The values are mean ± SD for 5 rats, *p<0.05 (one-way ANOVA followed by Bonferroni post hoc test).
Acknowledgments

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Abbreviations

α-Toc, α-tocopherol; γ-Toc, γ-tocopherol; Toc3, tocotrienol; TG, triacylglycerol; T-Chol, total cholesterol.

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