Antioxidant activity of vitamin B₆ delays homocysteine-induced atherosclerosis in rats

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Elevated plasma homocysteine is a risk factor for atherosclerotic disease. In the present study, we have examined whether the oxidative stress due to a low level of vitamin B₆ accelerates the development of homocysteine-induced atherosclerosis in rats. First, the effect of homocysteine thiolactone intake (50 mg/kg per d) on vascular integrity, lipid peroxide concentration, endothelial NO synthase (eNOS) expression and biochemical profiles was examined at day 1, day 21 and day 42 (five rats per group). The histochemical staining of the rat aorta showed no change at day 1 and day 21, but the subendothelial space was observed to be enlarged in rat aorta at day 42 with exposure to homocysteine thiolactone. Expression of eNOS was observed in rat aorta at day 42, but not at day 1 and day 21. Serum lipid peroxide concentration and biochemical profiles including glucose cholesterol and triacylglycerol showed no change at any day. Second, the effect of homocysteine thiolactone intake in the presence and absence of vitamin B₆ on vascular integrity was examined at day 1 and day 14 (five rats per group). Aortic lesions were observed in vitamin B₆-deficient rat aorta at day 14 but not in vitamin B₆-supplemented rats. The expression of eNOS was also observed in vitamin B₆-deficient rat aorta at day 14. Serum lipid concentrations of the vitamin B₆-deficient group significantly increased compared with concentrations of the vitamin B₆-supplemented group, though serum concentration of homocysteine did not change between both groups. These results suggest that the oxidative stress caused by a low level of vitamin B₆ accelerates the development of homocysteine-induced atherosclerosis in rats.

Antioxidants: Atherosclerosis: Homocysteine: Vitamin B₆

Atherosclerosis is the major cause of heart attack and stroke, which constitute the most prevalent diseases (Ross, 1999). The first step of atherosclerosis involves endothelial dysfunction as proposed in ‘the response-to-injury hypothesis’ by Ross (1993). The factors causing endothelial dysfunction include shear stress, oxidised LDL and oxidative stress. When endothelial dysfunction occurs, the endothelial cells become active and secrete adhesion molecules, cytokines, chemokines and growth factors through induction of transcription factor NF-κB. Subsequently, the adhesion molecules, such as E-selectin, intercellular adhesion molecule-1 and vascular cell adhesion molecule-1 mediate leucocytes rolling and adhesion to the endothelium. The migration of leucocytes from the blood vessels to the endothelium is promoted by chemokines such as monocyte chemotactic protein-1. The intimal macrophages and smooth muscle cells which are stimulated and transformed by cytokines engulf oxidised LDL and form fatty streaks. Ultimately, these fatty streaks develop into fibrous plaques, which rupture and cause thrombosis that may lead to acute coronary syndrome and myocardial infarction (Hansson, 2001).

Though hypercholesterolaemia has been suggested to be an important risk factor for atherosclerosis, most of the patients with heart attack myocardial infarctions show normal serum concentrations of cholesterol. Another factor associated with the formation of atherosclerotic lesions is hyperhomocysteinaemia (McCully, 1969). Recently, many studies have correlated associated high plasma or serum concentration of homocysteine with an increased risk of atherosclerosis (Eikelboom et al. 1999; Hankey & Eikelboom, 1999; Brattstrom & Wilcken, 2000; Ueland et al. 2000; Hofmann et al. 2001). Hyperhomocysteinaemia is a risk factor also in men. There seems to be a strong synergistic effect between hyperhomocysteinaemia and all traditional vascular risk factors (Taylor et al. 1991; Graham et al. 1997; van den Bosch et al. 2003). Furthermore, plasma homocysteine is a strong
Vitamin B₆ delays atherosclerosis

predictor of both cardiovascular and non-cardiovascular mortality in a general population aged 65–72 years old (Vollset et al. 2001). As homocysteine is a sulphhydryl-containing amino acid, and is derived from the metabolism of dietary methionine, the reactive oxygen species generated during oxidation of homocysteine may damage endothelial cells and promote accumulation of platelets (Harker et al. 1976). Because vitamin B₁₂ and folate act as cofactors for the re-methylation of homocysteine and vitamin B₆ for trans-sulfation to cysteine, low levels of these vitamins may cause hyperhomocysteinemia and increase the risk for atherosclerosis (Verhoef et al. 1996, 1997; Siri et al. 1998). Interestingly, Robinson et al. (1995) reported that low pyridoxal ⁵-phosphate confers an independent risk for coronary artery disease. Actually, it has been reported that folate and vitamin B₆ levels may be important in the primary prevention of CHD (Rimm et al. 1998).

On the other hand, many determinants of plasma homocysteine include genetic, physiological, and lifestyle factors, various diseases, and drugs are suggested (Refsum et al. 2004). Many of these factors cause a change in homocysteine concentrations by altering the function or blood concentration of the B vitamins (Ubbink, 1994), in particular folate and cobalamin, and/or by influencing renal function or, more rarely, by influencing enzyme activities.

Vitamin B₆ is an essential water-soluble vitamin required for normal growth and development (Tryfiates, 1980). Its physiologically active form, pyridoxal ⁵-phosphate, is derived from inactive precursors and functions as a cofactor in numerous enzymic reactions of amino acid metabolism. Apart from its function as a cofactor, antioxidant activities of vitamin B₆ are reported (Ehrenshaft et al. 1999; Stocker et al. 2003; Chen & Xiong, 2005). Pyridoxine seems to quench singlet oxygen at a rate comparable with that of vitamins C and E, two of the most highly efficient biological antioxidants (Ehrenshaft et al. 1999). Furthermore, Ullegaddi et al. (2004) reported that B-vitamin supplementation may have antioxidant and anti-inflammatory effects in stroke disease independent of a homocysteine-lowering effect. Thus vitamin B₆ seems to be involved in active-oxygen resistance. Based on these reports it is proposed that the antioxidant activity of circulating vitamin B₆ will suppress the progression of homocysteine-induced atherosclerosis independently from the action of homocysteine itself.

In the present study, we have examined whether the oxidative stress due to a low level of vitamin B₆ accelerates the development of homocysteine-induced atherosclerosis in rats.

Materials and methods

Animals and diets

Young 3-week-old male rats (40–50 g) (Wistar; Kyudo, Kumamoto, Japan) were used. The control group (n 5) had free access to standard chow (CE-2; CLEA Inc., Tokyo, Japan) and tap water. The experimental group (n 5) was orally given homocysteine thiolactone (Sigma, St Louis, MO, USA) in tap water (50 mg/kg body weight per d) for 42 d (Fig. 1(A)). Furthermore, to examine whether a low level of vitamin B₆ affects atherosclerosis, 3-week-old male rats (40–50 g) were subjected to experiments as shown in Fig. 1(B). One group (n 10) was fed with control diets for 4 weeks, and another group (n 10) was fed with vitamin B₆-free diets (Oriental Yeast Co., Ltd, Tokyo, Japan) to produce the vitamin B₆-deficient rats. After 2 weeks of feeding, five rats of each group were given homocysteine thiolactone in distilled water and others were given only distilled water for 2 weeks. Rats were kept at room temperature (23°C) under 12 h light–dark cycles. Body weight, food intake and amount of drinking water were recorded every other day. Rats were killed at the end of the experiments and liver was taken for the examination of glutamate–oxaloacetic transaminase and glutamate–pyruvate transaminase activities (units/mg protein). Blood was taken from the abdominal aorta and the serum was collected. The aorta was removed for histological studies. The present study was approved by the Kagoshima University Animal Committee (Japan) and the rats were maintained in accordance with the Kagoshima University guidelines for the care and use of laboratory animals.

Biochemical analysis

Serum biochemical data, glucose, triacylglycerol and total cholesterol were analysed to examine the nutritional condition of experimental animals. The serum concentration of triacylglycerol was estimated by a triglyceride G-test Wako (Wako Pure Chemical Industries, Ltd, Osaka, Japan). The serum glucose and total cholesterol were determined using SPOT-CHEM™ (Arkray Inc., Kyoto, Japan). The activities of glutamate–oxaloacetic transaminase and glutamate–pyruvate transaminase of liver tissue were analysed using the methods.
of Okada & Ochi (1971). Total protein was measured with BCA protein assay reagent (Pierce Biotechnology Inc., Rockford, IL, USA). The serum level of homocysteine was measured by homocysteine microplate (Substrate Trapping Enzyme) assay kit (Diazyme Laboratories, San Diego, CA, USA). The serum lipid peroxide concentration was determined by detection of thiobarbituric acid-reactive substances (TBARS). Thiobarbituric acid reagent (1 ml) was added to a tube containing 20 µl serum and 10 µl of 5 % butylated hydroxytoluene in ethanol. The tube was vortexed and heated at 100°C for 45 min. After centrifugation at 650 g for 10 min, the supernatant fraction was filtered (Millex-HV, 0.45 µm; Nihon Millipore Ltd, Tokyo, Japan) and the concentration of thiobarbituric acid-reactive substances was examined using a HPLC LC-10AD (Chromatopac, Shimadzu, Kyoto, Japan) attached C18 ODS column. The wavelength used for excitation was 515 nm and for emission 553 nm.

**Immunohistochemistry for endothelial nitric oxide synthase**

The tissue was deparaffinised (xylene 3 ×, 100 % ethanol 2 ×, 90 % ethanol, 80 % ethanol, 70 % ethanol, deionised water 3 ×) and heated in a microwave oven at 500 W for 15 min in a 10 mM-sodium citrate buffer (pH 6.0). After cooling the slides in the buffer for 20 min, the tissue was washed, incubated with 0.3 % H2O2 in 100 % methanol for 30 min. After washing with PBS (pH 7.4), the tissue was incubated with a blocking solution of 1.5 % normal goat serum for 1 h at room temperature. Then the slides were incubated with polyclonal antibody to endothelial nitric oxide synthase (endothelial NO synthase (eNOS)) (2 µg/ml; Affinity BioReagents Inc., Golden, CO, USA) overnight at 4°C. After washing with PBS, biotinylated goat anti-rabbit immunoglobulins (Dako Japan Co., Ltd, Kyoto, Japan) were applied and incubated for 30 min at room temperature. The presence of eNOS was detected using the Vectastain ABC kit (Vector Laboratories, Inc., Burlingame, CA, USA). Then the sections were visualised with 3,3′-diaminobenzidine tetrahydrochloride containing 0.05 % H2O2. Finally, the sections were washed, counterstained, dehydrated, cleared in xylene and mounted.

**Statistical analysis**

Data are presented as mean values and standard deviations. The statistical analysis was used Student’s t test for comparison of the subendothelial region and eNOS-positive region, and the Kruskall–Wallis test was used for comparison of the concentration of homocysteine and thiobarbituric acid-reactive substances. Analyses were performed using STATISTICA 4.5 software (Statsoft Japan Inc., Tokyo, Japan) on a Windows platform.

**Results**

Rats treated with homocysteine for 42 d showed the same growth curve as control rats. Also the biochemical data of serum such as glucose, triacylglycerol and cholesterol were almost similar in the two groups (Table 1). On the other hand, the histochemical staining of endothelium showed that subendothelial space enlargement increased in the homocysteine group after 42 d, probably due to the accumulation of lipid (Fig. 2 (A)), but the aorta from control rats did not show any change (P<0.001; Student’s t test; Fig. 2 (B)). In addition, the immunohistochemical analysis for eNOS showed the positive staining in the endothelial cells of B6-deficient rats given homocysteine. The expression of eNOS increased in the aorta from rats treated with homocysteine but not in control rats (Fig. 2 (C)).

To study vitamin B6-mediated vascular effects in vivo, rats were fed B6-deficient diets. After 2 weeks of the feeding, five rats of each group were given homocysteine thiolactone in distilled water and the others were given only distilled water for 2 weeks. The body weight of B6-deficient rats decreased from 2 weeks after feeding B6-deficient diets, characteristic of B6-deficient rats (body weight of B6-supplemented rats, 220.5 (SD 12.8) g; B6-deficient rats, 175.8 (SD 15.4) g) as suggested by Coburn et al. (1981). Moreover, glutamate–oxaloacetic transaminase and glutamate–pyruvate transaminase activities in rat liver, which are the B6 enzymes, decreased significantly in the liver of B6-deficient rats (glutamate–oxaloacetic

**Table 1. Serum concentrations of glucose, triacylglycerol (TG) and cholesterol in homocysteine (Hcy)-treated and control rats**

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<tr>
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<th>Day 21</th>
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<th>Day 42</th>
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<tbody>
<tr>
<td></td>
<td>Control</td>
<td>Hcy</td>
<td>Control</td>
<td>Hcy</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Glucose (mg/l)</td>
<td>860</td>
<td>52</td>
<td>750</td>
<td>102</td>
</tr>
<tr>
<td>TG (mg/l)</td>
<td>1480</td>
<td>102</td>
<td>1460</td>
<td>187</td>
</tr>
<tr>
<td>Cholesterol (mg/1)</td>
<td>386</td>
<td>92</td>
<td>428</td>
<td>116</td>
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transaminase activity was 13.8 (SD 1.2) for control liver and 8.7 (SD 0.7) for B6-deficient liver; glutamate–pyruvate transaminase activity was 4.1 (SD 1.1) for control liver and 1.43 (SD 0.6) for B6-deficient liver). Also, the serum concentrations of glucose, triacylglycerol and cholesterol did not appear to differ significantly among the four groups (Table 2).

The histochemical staining of the aortic tissue from B6-deficient rats treated with homocysteine showed that the subendothelial space increased after 14 d, whereas B6-deficient rats treated with only distilled water or control rats treated with distilled water or homocysteine showed no change (P < 0.001; Student's t test; Fig. 3 (A) and (B)). Furthermore, the immunohistochemical detection for eNOS showed the positive staining in the endothelial cells of B6-deficient rats given homocysteine (Fig. 3 (C)). B6-deficient rats treated with homocysteine showed the appearance of atherosclerosis-like alterations after 14 d, which was earlier than the period of 42 d when the atherosclerosis-like alterations appeared in the homocysteine-treated rats.

There was no statistical difference in serum homocysteine level among the B6-supplemented groups and the B6-deficient groups, though the level of serum homocysteine was higher in homocysteine-treated rats (P < 0.05; Kruskall–Wallis test; Fig. 4). However, the serum lipid peroxide levels in B6-deficient and homocysteine-treated rats were higher compared with those of control rats (P < 0.05; Kruskall–Wallis test; Fig. 5). These results suggest the earlier expression of endothelial dysfunction due to high levels of lipid peroxide caused by vitamin B6 deficiency.

Discussion

Homocysteine has been reported as an important risk factor of atherosclerosis (McCully, 1969; Hankey & Eikelboom, 1999; Hofmann et al. 2001; Adachi et al. 2002). Its oxidation generates reactive oxygen species (Loscalzo, 1996), increases cytokine production (van Aken et al. 2000), up-regulates adhesion molecules (Silverman et al. 2002), and induces endothelial dysfunction (Chambers et al. 1999). However, the mechanism by which homocysteine induces atherosclerosis remains unclear.

On the other hand, the concentration of homocysteine is regulated by the B vitamins B6, B12 and folate. Since vitamin B12 and folate act as cofactors for the re-methylation of homocysteine and vitamin B6 for trans-sulfation to cysteine, low levels of these vitamins may cause hyperhomocysteinaemia.

Table 2. Serum concentrations of glucose, triacylglycerol (TG) and cholesterol in B6-supplemented and -deficient rats

<table>
<thead>
<tr>
<th></th>
<th>B6-supplemented</th>
<th>B6-deficient</th>
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<tr>
<td></td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Glucose (mg/l)</td>
<td>860</td>
<td>62</td>
</tr>
<tr>
<td>TG (mg/l)</td>
<td>1524</td>
<td>186</td>
</tr>
<tr>
<td>Cholesterol (mg/l)</td>
<td>428</td>
<td>64</td>
</tr>
</tbody>
</table>

Hcy, homocysteine.
However, as vitamin B₆ has antioxidant activity apart from its role as coenzyme, the antioxidant activity of vitamin B₆ may suppress the homocysteine-induced atherosclerosis independent of homocysteine action itself.

In the present study, we examined the effect of the antioxidant activity of vitamin B₆ on the occurrence of hyperhomocysteinemia. Homocysteine-treated rats showed the structural alteration of endothelium after 42 d, whereas B₆-deficient rats treated with homocysteine showed the same effect after just 14 d. The level of serum homocysteine did not show any change but the level of serum lipid peroxide was increased in B₆-deficient rats. Serum homocysteine levels did not appear to change within a period of 14 d. Miller et al. (1992) reported that there was no difference in homocysteine concentrations between deficient and control animals after 6 weeks of the dietary regimen. These results suggest that the oxidant stress due to B₆ deficiency accelerates atherosclerosis. Indeed, Ehrenshaft et al. (1999) reported that pyridoxine quenches singlet oxygen at a rate comparable with that of vitamins C and E, two of the most highly efficient biological antioxidants. Interestingly, the findings of Bellamy et al. (1999) could be explained by folate itself having an antioxidant effect that opposes homocysteine-induced oxidative changes.

Furthermore, Barua et al. (2003) have reported that reactive oxygen species induce the up regulation of eNOS in vitro. The increase of eNOS expression observed in the present study is consistent with their findings and may support the decrease of antioxidant activity in B₆-deficient rats.

In conclusion, B₆ deficiency induced the oxidant stress which accelerates atherosclerosis and the antioxidant activity of vitamin B₆ appears to suppress homocysteine-induced atherosclerosis.
References


