Note

Effect of Vitamin E and C Supplements on Antioxidant Defense System in Cardiovascular Disease Patients in Zahedan, Southeast Iran

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Summary Oxidative stress plays an important role in the pathogenesis of cardiovascular disease (CVD). Growing evidence suggest that antioxidant vitamins might reduce the risk of disease outcomes by their ability to scavenge free radicals. The aim of the present study was to evaluate the supplementation of vitamins E and C on oxidant and antioxidant status in CVD patients. We conducted a case-control study with vitamin E (400 IU/d) and vitamin C (500 mg/d) supplementation in 40 CVD patients for 2 mo. Antioxidant (enzymatic and non-enzymatic) and oxidant status were analyzed pre and post supplementation. In the initial stage the activity of both enzymatic and non-enzymatic antioxidants were lower, while the malondialdehyde (MDA) level was elevated (p<0.0001). After intervention, a significant increase in superoxide dismutase (SOD) activity (61.7%), glutathione peroxidase (GPx) activity (59.3%), the levels of vitamin E (83.7%), C (145.3%), total antioxidant capacity (TAC) (62.8%) and a significant decrease in MDA (40%) value were observed (p<0.0001). There was a significant negative correlation between MDA and TAC. The results suggest that supplementation with a combination of vitamins E and C reduced lipid peroxidation and strengthened the antioxidant defense system. Hence, there will be beneficial effects on the heart by reducing oxidative stress in CVD patients.

Key Words vitamin E and C supplementation, lipid peroxidation, cardiovascular disease (CVD)

Oxidative stress plays an important role in the pathogenesis of cardiovascular disease (CVD). Numerous studies over the past decade have focused on the role of oxidative stress in CVD (1, 2). The effect on cardiovascular function may occur in two ways; either in the long-term development of atherosclerosis or in immediate damage during a heart attack or stroke (3). Both enzymatic—such as superoxide dismutase (SOD), catalase, and glutathione peroxidase (GPx)—and non-enzymatic antioxidants strengthen the system of the body. Lipid peroxidation, oxidation of low density lipoprotein (LDL) in particular, is the main causative factor in the development of CVD (4). During oxidation, peroxides followed by proxy radicals are formed which further degrade to aldehydes; malondialdehyde (MDA), hexanal and 4-hydroxynonenal (5). Antioxidant nutrients such as vitamins C and E are capable of inhibiting lipid peroxidation and reducing cardiovascular events (1, 6, 7). There is substantial evidence that free radicals affect the etiology of many diseases including vascular disease (8). At the same time the living organisms have antioxidant defense systems which prevent the damaging effects of excessive endogenously and exogenously produced free radicals (9). Non-enzymatic detoxification against reactive oxygen species (ROS) is determined by lipid-soluble antioxidants including vitamin E and carotenoids, and vitamin C, a water-soluble antioxidant present in the extracellular fluid of the arterial wall (10–12). Observational studies have concluded that high intakes of antioxidant vitamins such as vitamin E and C from food and supplements are related to low risk of CVD (13, 14). Some clinical trials have reported conflicting conclusions regarding the benefit of antioxidant vitamin supplement, which have not been confirmed (7, 9). Some studies have shown that mild to moderate deficiencies of vitamins E and C cause classic deficiency diseases, which may be involved in the development of CVD (11). So it is thought that antioxidant supplementation may help reduce the progression of CVD (1). Individuals with high dietary intakes of vitamin C, E and β-carotene have a lower risk of CAD (12). Many of the ingredients in commercial supplement packs have been demonstrated singly to have direct or indirect effect on the cellular antioxidant system (13).

Overall reported studies have shown inconsistent
results: beneficial effect of individual antioxidants, combined effect and no effect of vitamin E and C (1, 3, 8–10). Thus, the current evidence is insufficient to conclude that antioxidant vitamin supplementation reduces oxidative damage in humans. It will be important to clarify the discrepancy between the results of different studies. The aim of this study was to examine the combined effect of supplementation of vitamins E and C on oxidant and antioxidant defense systems in Iranian CVD patients.

Materials and Methods

Subject characteristics. Forty patients with CVD (57.7±10.6 y) were enrolled in the study. The diagnosis of CVD patients, admitted to the intensive cardiac care unit (CCU and Post-CCU) wards at Khatam-Al-Anbia Hospital of Zahedan University of Medical Sciences, was established according to clinical criteria (unstable angina, and chest pain which lasted for up to 3 h), echocardiography changes and total creatine kinase (CK) and CKMB activity.

Sixty-three healthy subjects (56.4±11.3 y) were selected as the control group. They did not have any history of disease and were not on any medication. None of the study subjects received vitamin supplements in the 3 mo prior to the study. The protocol of the study was approved by the local ethical committee of Zahedan University of Medical Sciences and informed consent was obtained from all patients and healthy subjects.

Interventional study. The vitamin antioxidant supplementation was carried out by supplying both vitamin C (500 mg L-ascorbic acid, Darou Pakhsh Pharmaceutical Mfg. Co, Tehran, Iran) and vitamin E (400 IU, dl-a-tocopheryl acetate, manufactured by international agencies for Health Burst, Davie, FL, USA). The patients were instructed to consume vitamin C and vitamin E daily for the duration of 2 mo. After 2 mo of supplementation, the effect of supplements was assessed by measurement of oxidant and antioxidants markers in the patients.

Blood sampling and analysis. Following a 12-h fast period, venous blood samples were collected in different tubes, before and after intervention and once for the control group. Heparinized tubes were used for measurement of MDA, total antioxidant capacity (TAC), SOD, GPx, and vitamin C and tubes without anticoagulant were used for the measurement of vitamin E. Plasma and serum were obtained by centrifugation at 3,000 rpm for 10 min and stored at −80˚C until analyzed.

MDA assay. Plasma MDA level was measured based on the reaction between MDA and thiobarbituric acid (TBA) using the method described by Satoh (14). The absorbance was read at 530 nm and expressed as micromoles of malondialdehyde per liter.

TAC. Total antioxidant capacity of plasma was measured by the ferric reducing ability of plasma (FRAP) assay. At low pH, reduction of ferric-tripyrilidyltrazine complex to the ferrous form displays an intense blue color with an absorption maximum at 593 nm (15).

SOD and GPx activities. SOD and GPx were measured using commercially available kits supplied by Randox Laboratories Ltd (Randox, UK) as described previously (16).

Vitamins E and C assay. Serum levels of vitamin E were determined using high pressure liquid chromatography (HPLC) as described previously (16). Plasma concentration of vitamin C was measured by a spectrophotometric method (17) as described previously (16).

Statistical analysis. Results are expressed as means±SD. Statistical analysis was performed using SPSS 17 software (SPSS Inc, Chicago, USA). Paired and unpaired sample t-tests were used to compare the groups. p-values less than 0.05 were considered statistically significant.

Results

The demographic and clinical characteristics of CVD patients and healthy subjects are shown in Table 1. During the experimental period, 9 patients withdrew due to open heart surgery or transferred to another city for continuing treatment.

The levels of vitamin C, vitamin E, SOD, GPx, MDA, and TAC among normal subjects and CVD patients (before and after 2 mo supplementation) are shown in Fig. 1. After 2 mo of supplementation, plasma level of vitamin C increased significantly (almost 1.5 fold) (p<0.0001) from baseline and reached the normal level.
Similarly, the level of vitamin E increased significantly (83.7%) \( (p<0.0001) \) so that it was higher than in normal subjects \( (p<0.001) \) (Fig. 1B). Despite significant increase in the activities of SOD (61.7%) and GPx (59.3%) \( (p<0.0001) \), they were still significantly lower than for the control group \( (p<0.001) \) (Fig. 1C and D). Plasma levels of MDA considerably decreased (40%) \( (p<0.0001) \) from baseline and reached the same level as the control value (Fig. 1E). Plasma total antioxidant capacity (TAC) after the 2-mo experimental period, increased significantly (62.8%) \( (p<0.0001) \) and reached that of the healthy individuals (Fig. 1F). A negative significant correlation was observed among MDA with GPx \( (r=-0.37, \ p=0.04) \) and TAC \( (r=-0.30, \ p=0.003) \) after intervention.

**Discussion**

The role of free radicals in the development of complications of CVD has been established by various epidemiological and clinical trials with antioxidants \( (8, 10, 11, 13) \). The increased level of MDA and lipid peroxidation products confirms the oxidation of LDL in CVD subjects \( (18) \). After 2 mo of vitamin E and C supplementation, MDA levels fell to almost normal values. Though vitamin C deficiency has been shown to be a risk factor for CHD \( (7, 9, 11, 19, 20) \), some studies have indicated that there is an inverse association between CVD and high dietary vitamin C intakes or high plasma vitamin C concentrations, whereas other studies have not found any association between CVD and high plasma vitamin C concentration or use of supplements \( (9, 19) \). Vitamin C may enhance endothelial function by scavenging oxy-
showed that supplementation of the combination of a period of 2 mo in the present study, our results copherol acetate and 500 mg ascorbic acid was tried for did not have any effect (26). However, although supplementation with the combination dose of 400 IU α-tocopherol acetate and 500 mg ascorbic acid was tried for a period of 2 mo in the present study, our results showed that supplementation of the combination of vitamins E and C reduced lipid peroxidation, and both vitamins acted as free-radical scavengers. These vitamins also strengthened the antioxidant defense system against oxidative damage, which was confirmed by the inverse correlation among MDA with GPx and TAC. Hence, supplementation of vitamins E and C may have beneficial effects in CVD patients which remain to be clarified.

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Conflict of Interests: None.

REFERENCES


