

## Plasma level of vitamin C in women with pre-eclampsia in Ardabil, Iran

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### Abstract

**Background:** Pre-eclampsia, a vascular disorder of pregnancy, is a leading cause of maternal morbidity as well as perinatal morbidity and mortality. The cause of pre-eclampsia remains largely unknown. Accumulating evidences from clinical and epidemiological studies suggest that diffuse endothelial dysfunction, resulting from oxidative stress, plays an important role in the pathogenesis of pre-eclampsia. Independent effect of maternal biological markers, such as elevated triglyceride or low plasma vitamin C (ascorbic acid) concentrations are related to increased pre-eclampsia risk.

**Objectives:** The aim of this study was to determine plasma concentrations of vitamin C in women with pre-eclampsia.

**Materials and Methods:** This cross-sectional, case controlled study comprises 40 pre-eclamptic and 80 normo-tensive pregnant women of singleton gestations in third trimester, in Allavi Hospital, Ardabil, Iran. Blood samples (5ml) were collected aseptically in heparin tube, and the plasma concentration of vitamin C was determined by spectrophotometric method. SPSS software package was used for analyzing the data.

**Results:** Plasma vitamin C level was found to be significantly lower ( $p<0.001$ ) in the pre-eclampsia group than this level in the control group. Vitamin C levels in the pre-eclampsia group were found to be influenced by their maternal age ( $p<0.01$ ), and was positively related to the gestational age ( $p<0.01$ ).

**Conclusion:** Knowledge from this study may contribute to the development and evaluation of behavioral and medical interventions aimed at reducing the occurrence of pre-eclampsia.

**Key words:** Pre-eclampsia, Pregnancy, Vitamin C

### Introduction

Pre-eclampsia affects between 0.4% to 2.8% of all pregnancies in developed countries and even more in developing countries, leading to as many as 8,370,000 cases worldwide per year (1). Maternal mortality and morbidity from hypertensive disorders of pregnancy, including pre-eclampsia, remain high worldwide and represent a troublesome issue of modern perinatology (2). The risk of pre-eclampsia significantly increases in women with previous history of pre-eclampsia and in those with either preexisting vascular diseases or conditions associated with increased cardiovascular risk, such as renal diseases, hypertension, diabetes, thrombophilia, and obesity (body mass index  $>29$ ) (3). However, aetiology and pathogenesis of the pre-eclampsia is still unknown.

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Pathophysiological characteristics of pre-eclampsia contain altered vascular reactivity, loss of vascular integrity, activation of coagulation cascade and inadequate trophoblastic invasion of spiral arteries, which suggests dysfunction of the vascular endothelium (4-7). In addition, It has been documented that pre-eclampsia is accompanied by oxidative stress that contributes to vascular dysfunction (6-8). A recent study in which oxidative stress was investigated in maternal blood by a wide variety of markers showed evidence of oxidative stress in pre-eclampsia by some (notably reduced plasma vitamin C concentrations), but not all markers used, leading to the conclusion of a mild state of oxidative stress in blood of pre-eclamptic women (9).

The inadequate trophoblastic invasion results in a high resistance and low flow uteroplacental circulation that develop placental ischaemia and hypoxia (7). These could trigger lipid peroxidation and induce endothelial damage in pre-eclampsia (6,8).

Oxidative stress plays an important role in the pathophysiology of pre-eclampsia. It has been

postulated that the placental ischaemia initiates excessive free radical production and triggers multiple chain reactions, which ultimately results in widespread endothelial damage. Under hypoxic condition, the enzyme xanthine dehydrogenase is shifted to the oxidase generating free oxygen radicals (7) that attach unsaturated lipids and thiol containing proteins in cell membranes (10). This oxidative attack causes damage to enzymes, cell membranes, proteins and nuclear materials. The superoxide anion, an important free radical, can inactivate endothelium-derived-relaxing factor (11) and inhibit prostacyclin synthesis (12,13) resulting in the platelet aggregation and vasospasm, which are the two important characteristics of pre-eclampsia. Thus, oxidative stress arises from increased production of reactive oxygen species or deficiency in antioxidant nutrients. It has been suggested that deficiency in antioxidant vitamins would developed pre-eclampsia (6,14). Antioxidant vitamins, with the ability to stabilise highly reactive free radicals, act as the first line of defense against free radical attack and lipid peroxidation (15). Vitamin E, vitamin C and beta-carotene (pro-vitamin A) are known to be powerful antioxidants (15-17).

Vitamin E ( $\alpha$ -tocopherol) is the major lipid-soluble antioxidant that protects cells against lipid peroxidation. Vitamin C is a quencher of free radicals as well as singlet oxygen. It also regenerates vitamin E. Beta-carotene, by quenching singlet oxygen, also acts as a potential antioxidant. Antioxidant vitamins have been reported to have an important function in regulating blood pressure (18). Nitrous oxide (NO) is the most important endothelium dependent vasodilator and is highly susceptible to oxidative damage. The antioxidants vitamins, C and E, are able to inhibit formation of free radicals thereby inhibiting the oxidation of NO, and thus maintaining the vasodilator status of blood vessels. Vitamin C has been reported to have a direct acute effect on the inhibition of the constrictor response of resistance arteries to a variety of stimuli. It is reported that there is synergism between the actions of vitamin E and vitamin C (19-20). However, there is little information regarding the functions of antioxidant vitamins particularly plasma concentrations of vitamin C in pre-eclampsia especially in Iranian populations. Therefore, we tried to evaluate the plasma concentrations of vitamin C in pre-eclampsia and normo-tensive pregnant women in Ardabil province, Iran.

## Materials and Methods

### Study population

This study was designed as a cross sectional case control study and was conducted in the department of

Obstetrics and Gynecology, Ardabil Educational Hospital "Allavi", Ardabil, Iran. The biochemical analysis was carried out in the department of biochemistry.

The study included 40 pre-eclampsia and 80 normotensive pregnant women of singleton gestation in third trimester. The subjects were selected under defined criteria. Pre-eclamptic patients were at 28 to 42 weeks of singleton gestation, with one measurement of diastolic pressure of 110 mmHg or more, or two measurements of 90 mmHg or more on two consecutive occasions 6 hours or more apart, and urinary protein of 2+ or more (100mg/dl; dipstick reagent strip, Boehringer Mannheim, Germany) (21). The exclusion criteria included; previous history of hypertension and proteinuria before conception or before 20 weeks of gestation, any associated disorders, a history of antioxidant vitamin therapy during last one year, and smoking. The patients who developed convulsions were considered to be suffering from eclampsia and were excluded. As cohort control, age and socio-economically matched healthy normo-tensives women at 28 to 42 weeks of singleton gestation with no urinary protein were recruited by convenience.

### Body Mass Index (BMI)

BMI is a widely used method of calculating percentage of body fat as a continuum rather than simple conventional classification units. This study tries to determine the mechanisms by which body fat increases pre-eclampsia risk.

BMI was calculated as weight before pregnancy (kg) divided by squared height ( $m^2$ ).

### Plasma analysis

Peripheral blood samples (5ml) were collected from antecubital vein of cases and controls in heparin tubes. The samples were kept undisturbed on wet ice for at least 10 minutes, protected from ultraviolet light, and were processed within 30 minutes. The samples were centrifuged at 3000 rpm for 10 minutes in a cooling centrifuge. Plasma was decanted into cryovials, preserved with metaphosphoric acid/dithiothreitol solution, and stored at -24°C for analysis. The concentration of ascorbic acid in the plasma was determined by spectrophotometric within 48 hours as described by Kyaw (22). Absorbance was measured against a blank reagent at 700 nm by a spectrophotometer (UV-VIS, Ependorf Co., Germany). Plasma vitamin C was categorized as deficiency (less than 0.2 mg/dl), marginal deficiency (0.2-0.4 mg/dl) and normal (more than 0.4 mg/dl).

### Statistical analysis

SPSS software package (version 10.0, SPSS Inc. Chicago, USA) was used to analyse the data.

Descriptive statistics were calculated for all variables. The associations of plasma vitamin levels and BMI were assessed using multiple regression analysis. In the regression model, plasma vitamin levels, maternal age and body mass index (BMI) were adjusted. The results were statistically analyzed using student's t-test and regression analysis. Values were expressed as percentage and mean  $\pm$  SE.

## Results

Clinical parameters of the patients and normotensive controls are shown in Table I. The mean maternal and gestational ages of the patients and controls were similar. Patients had different gravida distribution and had nearly equal proteinurea. The systolic and diastolic pressures of pre-eclamptic patients were significantly ( $p<0.001$ ) higher than controls (Table I).

**Table I.** Maternal Age and clinical characteristics of pre-eclamptic and normo-tensive pregnant women

Clinical characteristic	Normo-tensive	Pre-eclamptic
Maternal age (year)	22.05 $\pm$ 1.41	27.11 $\pm$ 0.93
Gestational age (week) at sampling	34.64 $\pm$ 0.95	37.23 $\pm$ 1.64
Gravida in no (%)		
primi	12 (30)	28(35)
multi	7(17.5)	12(15)
Nulliparous	21(52.5)	40(50)
Proteinurea		
primi	0	2+(26) <sup>1</sup>
multi	0	3+(18)
BMI (kg/m <sup>2</sup> )	28.5 $\pm$ 0.4	35.7 $\pm$ 0.6 <sup>2</sup>
Systolic blood pressure (mmHg)	109.86 $\pm$ 9.27	160.68 $\pm$ 22.61 <sup>3</sup>
Diastolic blood pressure (mmHg)	73.43 $\pm$ 8.81	109.16 $\pm$ 15.21

Values are expressed as mean  $\pm$  SE

Compare Means: Independent-samples t-test

1,2  $p<0.001$

3  $p<0.01$

Odds ratio (OR) 2.1 (95% confidence interval: 1.4, 3.4)

Table II shows plasma vitamin C concentration in the cases and controls. It was observed that vitamin C concentration in pre-eclamptic patients was significantly lower than control ( $p<0.001$ ).

Vitamin C level in pre-eclamptic patients was found to be influenced by maternal age ( $p<0.01$ ). In total, the correlation between the age and vitamin C level was determined and it was found to be positively related to each other ( $p<0.01$ ). In controls, gestational age showed positive influence on vitamin C level ( $p<0.01$ ).

**Table II.** Serum concentrations of Vitamin C in pre-eclampsia and normotensive pregnant women

Antioxidant Vitamin (mg/dl)	Normo-tensive		Pre-eclampsia	
	n	mean $\pm$ SE	n	mean $\pm$ SE
Vitamin C $\leq$ 0.2	3	0.34 $\pm$ 0.11	39	0.11 $\pm$ .30*
Vitamin C $>$ 0.2	77		1	

Values are expressed as mean  $\pm$  SE

\*  $p<0.001$

## Discussion

The etiology of pre-eclampsia is elusive. There is no definite way to prevent pre-eclampsia. Oxidative stress is thought to be associated with pre-eclampsia (5-7). From early pregnancy, the human placenta influences maternal homeostasis. It is rich in mitochondria and when fully developed consumes about 1% of the basal metabolic rate of the pregnant woman (23). Placenta is also highly vascular and is exposed to high maternal oxygen partial pressure. These characteristics explain, in part, the generation of superoxide, because about 5% of all electrons in the mitochondrial respiratory chain leak out of the mitochondrion (24,25). Initially the placenta, has a hypoxic environment. As it matures and its vascularization develops, it changes to an oxygen-rich environment and its abundant mitochondrial mass favors the production of reactive oxygen species (ROS) (26).

Vitamin C deficiency affects placental structure, ROS production and facilitates placental infection, which result in increased risk of premature rupture of placental membranes and premature births (27,28). The placenta is avid to absorb vitamin C so when maternal plasma ascorbic acid concentration is low it is absorbed by active mechanisms. At higher plasma ascorbic acid concentrations, it enters the placenta by passive diffusion. Curiously, ascorbate is preferentially taken up in the oxidized form (dehydroascorbic acid), which more easily passes the lipid layer. However, it is transformed to its reduced form at the expense of other reducing agents, and donate electrons before transferring to the fetus by active transport (29).

Plasma antioxidant activity has been reported to increase progressively throughout pregnancy (4,10,18). This study investigated plasma vitamin C concentration in pre-eclampsia and attempted to evaluate its correlation with the etiology of this disease.

Some reports documented an increase in plasma vitamin E levels in pre-eclampsia (10,27), but others have found a decreased concentration (5,7).

Vitamins E (alpha-tocopherol) and C, have differences in the contribution they make to antioxidant potential, as vitamin E is the major lipid-soluble chain-breaking antioxidant in cell-membranes while vitamin C is an important aqueous phase antioxidant.

Antioxidants may act synergistically, for instance when vitamin C regenerates alpha-tocopherol from the tocopherol radical (30), this 'sacrificial' antioxidant acts more by sparing vitamin E than by recycling (31). Thus, it might be important to evaluate the effectiveness of potential antioxidant defense systems in limiting scale.

This study provides evidence for the relationship between plasma vitamin C levels during the third trimester and pre-eclampsia. Our results indicated the plasma vitamin C concentration to be significantly lower in pre-eclampsia, as compared to control group which is similar to the other studies (32-35).

Ascorbic acid has been described as the major 'front line' water-soluble antioxidant. Vitamin C can regenerate vitamin E from the alpha-tocopheroyl radical. In addition, it has also been documented that vitamin C, as an effective antioxidant in human plasma, provides major defense against the diseases caused by oxidative stress (18,19).

It is likely that severe oxidative stress in eclampsia utilises a higher amount of vitamin C to fight the oxidative stress leading to a depletion of the vitamin. The higher vitamin C levels in pre-eclampsia may serve to prevent oxidation of NO (endothelium dependent vasodilator) to maintain the vasodilatation of blood vessels (18).

The vitamin C levels in pre-eclampsia showed positive correlation with maternal age which is in agreement to some extent with the report of Kharb (35).

Deleterious effects of free radicals include initiation of lipid peroxidation, oxidative damage of biomolecules, and cellular dysfunction, which may initiate maternal vascular endothelial dysfunction and leukocyte activation (20). Free radical chain oxidation and the interaction of various antioxidants are now attracting the attention of nutritionists (32). The important role of vitamin C in pre-eclampsia, suggests that changes in its concentration may influence susceptibility of vascular endothelium to oxygen toxicity (36). Thus, analyzing vitamin C concentration may provide a means of assessing the total capacity of the chain-breaking antioxidants to prevent lipid peroxidation in plasma.

## Conclusion

The results of our study showed that plasma vitamin C concentration of the pre-eclamptic women was significantly lower than that of controls. Vitamin C level in pre-eclampsia was also found to be influenced by maternal age, gestational age and BMI. More studies should be done before pre-eclampsia is clinically evident. Vitamin C assessment should be done at different stages of pregnancy with tools validated for each stage of pregnancy.

In addition, in view of low plasma vitamin C concentrations in pre-eclampsia, a combination of vitamins C and E is a promising prophylactic strategy for prevention of pre-eclampsia.

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