

## Research Article

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## Serum levels of vitamin D, calcium, phosphorus, and oxidative parameters in healthy and diabetic people

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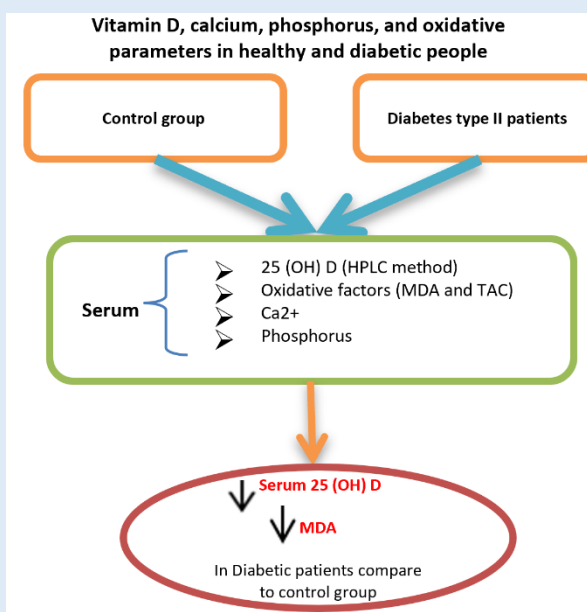
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### ABSTRACT

**Introduction:** Diabetes mellitus is a metabolic disease that is a primary public health consideration. Low Vitamin D levels are linked to type 2 diabetes (T2DM), diminished insulin release, and enhanced insulin resistance in humans and animals. Vitamin D is also involved in the regulation of calcium and phosphorus homeostasis. Oxidative stress and antioxidant imbalances are important for the progression of diabetes as well. In this endeavor, the levels of vitamin D, calcium, phosphorus, and evaluation of the oxidant-antioxidant factors of malondialdehyde (MDA) and total antioxidant capacity (TAC) in healthy and diabetic people were compared.

**Methods:** This descriptive-analytical study was conducted in 2020 in Shiraz, Fars province, Iran. The population included 40 T2DM patients (with HbA1c equivalent 6-8) without comorbidities, 20-60 years old for both genders,



and 40 healthy individuals (female and male between 20-60 years old without comorbidities). The high-performance liquid chromatography (HPLC) method was adopted for measuring Vitamin D and for measuring other levels, the colorimetric method was used. Using SPSS 22, statistical analysis was performed. The Mann-Whitney U test for quantitative data was applied.  $P < 0.05$  was deemed significant.

**Results:** There was a statistically significant difference between the two groups when it came to the means of vitamin D and MDA. In the diabetic group, vitamin D levels were lower ( $p = 0.001$ ) and MDA levels were higher ( $p < 0.001$ ). Comparing the level of calcium and phosphorus in diabetics and healthy people revealed no significant difference. This result was also true for the TAC test.

**Conclusions:** According to our results, the mean of vitamin D in T2DM was significantly lower than healthy people and MDA in T2DM significantly increased compared to the control group, suggesting that increasing the activity of this enzyme in the development of secondary complications in diabetic patients is a predisposing factor.

**Keywords:** Vitamin D, Diabetes mellitus, HPLC, Oxidative stress

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## INTRODUCTION

Type 2 diabetes mellitus (T2DM) is characterized by a chronic rise in blood sugar due to a relative lack of insulin, insulin resistance, or both. Diabetes and its late effects lead to reduced life expectancy and health costs [1]. T2DM has become one of the most momentous and common health problems in recent decades and is now the 7th leading cause of death in the United States as well as worldwide with 5.2 million deaths. Diabetes mortality is estimated at 82.4 per 100,000 [2].

T2DM is a complex metabolic disease that causes a range of insulin resistance in the target tissues or defects in insulin secretion from the pancreas [3]. Various predisposing factors for this disease have been mentioned including systemic inflammation, oxidative stress, obesity, lifestyle, diet, and the role of vitamins and minerals [3, 4].

During the last decade, vitamin D has been linked to an increased risk of T2DM, and vitamin D

supplements are thought to be an effective option in lowering the risk of T2DM. Recently, various studies have been performed to determine the effect of vitamin D supplementation on the control of diabetes in T2DM [5-7]. Previous studies suggested that receptors of Vitamin D are located within pancreatic beta cells, and adequate levels of vitamin D aids insulin sensitivity and insulin secretion [8, 9]. Moreover, vitamin D is involved in calcium regulation and through this role, it can indirectly affect insulin release from pancreatic beta cells [1].

Contradictory results have been reported in previous studies regarding the effect of vitamin D on metabolic syndrome and related diseases. For combined vitamin D and calcium supplements, intervention trials have only been effective in high-risk populations (eg. those with glucose intolerance) and have reduced the incidence of T2DM [10]. However, in RCT studies, when the study was faced with low sample size, an inappropriate dose of vitamin D, low homogeneity of vitamin D status in the

samples, and genetic polymorphism (at least for vitamin D receptor), the results have shown that vitamin D and calcium do not play a protective role in T2DM [11]. Vitamin D receptors are scattered in the pancreatic beta cells and immune system cells. In addition, vitamin D is involved in the activity of endopeptidases in calcium-dependent beta cells and can both directly induce beta cells to secrete insulin and by indirectly increasing intracellular calcium, convert calcium-dependent channels in beta cells which convert pro-insulin to insulin [12].

Hyperglycemia also causes oxidative stress by increasing protein glycation and glucose oxidation. Overproduction of reactive oxygen species and consequent depletion of the body's antioxidant defense system leads to lipid peroxidation and cell damage which then lead to macro-and microvascular complications of diabetes [13].

Due to the importance of vitamin D deficiency and macro-mineral deficiency in diabetic patients, differences in study results, differences in climatic conditions and diet in different regions, and the lack of similar studies in recent years in Shiraz city, this study aims to compare vitamin D levels, calcium, phosphorus, and oxidative conditions in diabetic and healthy patients.

## SUBJECTS AND METHODS

This descriptive-analytical study was conducted in the Laboratory of Pathobiology and Genetics of Peyvand Shiraz city, Fars province, south of Iran (2020). Approval was obtained from the ethics committee of the Shiraz University of Medical Science. The procedures used in this study adhere to the tenets of the Declaration of Helsinki.

**Inclusion and exclusion criteria:** The population of our study was selected randomly from diabetic patients who were referred to one of the medical diagnostic labs in Shiraz city. Inclusion criteria in our investigation were diabetic patients (T2DM) who

have been diagnosed for at least three years and with HbA1c equivalent to 6-8, those who are taking oral medicine for the control of blood glucose (other than insulin), 20-60 years old in both genders, non-smoker, non-alcoholic, and without comorbidities. Exclusion criteria included consuming supplements in the form of minerals and vitamin D, obesity (BMI>33), nephropathies, hepatitis, or any comorbidity that requires medical attention, those with malabsorption, infertility, oligomenorrhea, pregnancy, lactation, malignancy, use of drugs affecting bone metabolism (estrogen, calcitonin, bisphosphonates, etc.), and injection of vitamin D in the last six months.

**Collection of samples:** Forty healthy individuals were randomly selected from patients with no history of diabetes. The control group was matched to the patient group in terms of number, age, and sex.

The study process was explained verbally to all participants. After obtaining the informed consent of the study participants, study-related interviews were conducted with them. Those who wished to participate in this study used their blood serum to measure the amount of calcium, phosphorus, vitamin D, Malondialdehyde (MDA), and their total antioxidant capacity (TAC).

Sampling was performed after 8-12 hours of fasting from 8-11 a.m. and before taking blood glucose-lowering medications. Serum and plasma (containing the anticoagulant EDTA) were isolated to measure the target assays.

### **Estimation of Vitamin D, Calcium, Phosphorous, Malondialdehyde and total antioxidant capacity:**

Vitamin D was measured by high-pressure liquid chromatography (HPLC) according to the Galunská et al. method [14]. 0.5 ml of the sample were mixed with 1 ml of ethanol 96 (for 1 minute) to precipitate the protein. Then, the mixture was centrifuged for 10

minutes at 5500 rpm to achieve a clear supernatant. Subsequently, the clear liquid with 3 ml of hexane was re-extracted and evaporated under a slow flow of nitrogen. The dried residue was dissolved in 0.2 ml of methanol and HPLC analysis was carried out (*Waters pump binary 1525, USA*). Ten microliters of the prepared sample were injected into 25 cm long, 5-micron diameter columns, and vitamin D was measured by UV-visible detector [14].

MDA content, which determined the extent of lipid peroxidation quantitatively, was assessed using a colorimetric method according to the method prepared by the Zellbio (Germany) kit. Lipid peroxidation is measured by the reaction of thiobarbituric acid with MDA.

The TAC was measured using a colorimetric approach based on Ferric Reducing Potential of Plasma (FRAP).

Calcium and phosphorus were measured by the photometric method (*Pars Azmoon* diagnostic kits) and with an auto-analyzer (*BT-1500 Biotenica, Italy*).

**Statistical analysis:** Demographic and laboratory information were collected after coding and entered into the SPSS 22 software. The Shapiro Wilks test was used to evaluate the normality of the data. To compare the mean of the variables in the two groups,

the t-test was used if the data was normal and, if it was not normal, the Mann-Whitney test was used.

## RESULTS

**Demographics result:** The Mean±SD of the participant's age was 45±9 years (range from 20 to 60 years). There was an equal proportion of both sexes in the groups.

**Vitamin D result:** For vitamin D levels, there was a statistically significant difference between the two groups. The level of vitamin D in the diabetic group was 29.1 ±2.25 IU/mL and in the control group, it was 33.5 ± 4.5 IU/mL (p = 0.001) (Table 1).

**Calcium and phosphorus result:** Comparing the level of calcium and phosphorus in diabetics and healthy people revealed no significant difference (Table 1).

**Malondialdehyde and total antioxidant capacity results:** There was a significant difference between the two groups in terms of MDA level (p<0.001), and its value in the diabetic group was higher than the control. However, in the case of TAC, there was no significant difference between the two groups (Table 1).

**Table 1.** Serum levels of different variables in the two groups.

Variable	Diabetic (type 2)	Non-Diabetic	P-value
	Mean±SD	Mean±SD	
<b>25 (OH) Vit D</b> IU/mL	29.1± 2.25	33.5±4.5	<b>0.001</b>
<b>Calcium</b> mg/dL	9.53±0.2	9.64±0.18	<b>0.126</b>
<b>Phosphorous</b> mg/dL	3.62±0.37	3.47±0.28	<b>0.234</b>
<b>MDA</b> µM	2.23±0.28	1.08±0.19	<b>&lt;0.001</b>
<b>TAC</b> mM	1.57±0.41	1.66±0.23	<b>0.473</b>

MDA Malondialdehyde; TAC: Total antioxidant capacity; Vit D: Vitamin D; SD: standard deviation

## DISCUSSION

This study aimed to appraise the level of vitamin D, calcium, phosphorus, and oxidative status in diabetic and non-diabetic patients.

After conducting this study, we detected that the level of vitamin D in the T2DM group was significantly lower compared to the healthy group. It can be concluded that there is a role for vitamin D in regulating blood sugar and improving insulin sensitivity by reducing insulin resistance [1]. This study correlates well with the study conducted by Chiu *et al.* among 126 healthy and glucose tolerant subjects where 47 subjects were detected to have Vitamin D levels less than 20 ng/ml [8].

The most well-known use of vitamin D is for bone protection, but research has begun to explore its role in many areas of health. It has been proven that vitamin D deficiency can be a factor in developing Type 2 diabetes. It has also been demonstrated that the beta cells of the pancreas that secrete insulin contain the vitamin D receptor and the enzyme alpha 1 hydroxylase [15].

The study by Osmani was conducted on 50 newly diagnosed Type 2 diabetes mellitus patients for evaluating the levels of Vitamin D in diabetic and non-diabetic patients; they found that the diabetic group had lower levels of Vitamin D compared with the control. It can be concluded that there is a role for Vitamin D in regulating blood sugar levels and improving insulin sensitivity [1].

Moreira-Lucas *et al.* performed a double-blind, randomized, placebo-controlled 24-week trial to assess insulin sensitivity and beta-cell function. They used 71 participants whose serum vitamin D levels were  $25(\text{OH})\text{D} \leq 65$  nmol/L and had high fasting blood sugar and elevated glycosylated hemoglobin. For 24 weeks, once a week, cheese with and without vitamin D were given to subjects that were randomly divided

into two groups of 28,000 IU of vitamin D3 (VitD; n = 35) and placebo (n = 36). The results of their study after glucose tolerance testing in placebo and vitamin D groups showed that weekly doses of vitamin D3 in people with undesirable vitamin D levels and at risk for Type 2 diabetes could not improve oral glucose tolerance or other glycemic markers [16].

Calcium has been discovered to have an indirect effect on insulin secretion. Vitamin D can play a role in keeping extracellular calcium levels normal, thereby controlling the flow of calcium through the cell membrane. Decreased serum vitamin D levels can affect and reduce the ability of calcium to secrete insulin [1]. In our study, serum calcium and phosphorus levels did not show a statistical significant difference with the control group. In contrast to our study, Chen Hui *et al.* have shown that in Type 2 diabetic patients, the circulating phosphorus was lower while the circulating calcium was higher [17].

Reduced calcium absorption during vitamin D deficiency triggers the secondary release of parathyroid hormone and calcium reabsorption increases in the kidneys. This condition prevents the entry of calcium required for insulin processing in target cells and reduces insulin sensitivity by increasing intracellular calcium levels [18]. Decreased insulin sensitivity increases parathyroid hormone secretion [19]. Thus, increased parathyroid hormone due to low levels of vitamin D has an unfavorable effect on insulin release from beta cells [20-21].

Another way vitamin D affects diabetes is by increasing the expression of insulin receptors and improving insulin function, as well as increasing the insulin response to glucose transport [22]. Vitamin D prevents beta-cell apoptosis and thus maintains beta-cell mass [23].

Diabetes is often correlated with elevated oxidative stress as a result of decrease of antioxidant defenses and a rise in free radical generation [24]. Free radicals attack lipids to form malondialdehyde, one of the most important markers of lipid peroxidation, following an enzymatic reaction. Increased levels of malondialdehyde indicate a disorder of the antioxidant defense system [25]. Measurement of serum malondialdehyde levels is a diagnostic indicator of lipid peroxidation and oxidative stress, which is often examined in pathological conditions caused by free radicals [26]. The results of our study were similar to the Pieme et al. study regarding the serum level of malondialdehyde, which was significantly increased in T2DM compared with the control group [26]. Li et al., in 2019, indicated that the serum level of malondialdehyde in diabetics was significantly higher when compared to the healthy group [24]. Similar reports have been made in previous studies [27-28].

TAC represents the total peroxide damage caused by low molecular weight, enzymatic, free radical particles and reflects the effect of this damage on the enzymatic and non-enzymatic antioxidant balance in the body. Increased production of ROS in body tissues and fluids has been shown to reduce TAC [29]. Derived from the findings of the current research in T2DM and healthy individuals, the TAC of these individuals did not show a statistically significant difference. This result contradicts the findings of the study by Gunawardena et al. in 2019 and Pieme et al. 2017. In their study, the difference in serum levels of total antioxidants in T2DM compared to healthy individuals was reduced and this decrease was statistically significant [13, 26]. Also, in a study by Li et al., serum total antioxidant levels decreased in diabetics [24]. Since oxidative factors have reserves in the body, in patients with glycemic crises with

insufficient reserves of TAC tissue and reduced ability to produce TAC, we can see a decrease in this oxidative factor; if the body can reproduce it, they will not observe a significant change in the level of this factor [30].

Climatic conditions affect electrolyte levels, vitamin D, and blood glucose levels. Despite access to data related to these variables, few studies have been conducted in Shiraz. Some strengths of our study were that in selecting the sample, we applied a specific age limit and a specific duration of the treatment period, and the method of measuring vitamin D was done using the HPLC method. However, the use of the HPLC method and the cost of testing with this method limited the sample size.

## CONCLUSION

According to our results, the average level of vitamin D in diabetic patients was significantly lower than in healthy people. A lack of vitamin D can contribute to the long-term effects of diabetes, including cardiovascular disease. Due to the significant increase in MDA in T2DM patients compared to the healthy group, increased activity of MDA is a predisposing factor to the development of secondary difficulties in T2DM. By knowing more about the changes in the activity of antioxidant enzymes and oxidative factors in these patients in different stages of the disease and the factors affecting it, we can be more hopeful about increasing the effectiveness of pharmacological and nutritional interventions to reduce oxidative stress in diabetic patients. Fruits and vegetables have a protective role against chronic diseases such as cardiovascular disease, ocular and nerve diseases, stroke, cancer, diabetes, and high blood pressure. To prevent high blood pressure, stroke, cardiovascular disease, and other deficiencies related to micronutrients, it is recommended to consume at

least 400-500 grams of fruits and vegetables daily, because insufficient intake of these substances causes non-communicable nutritional disorders. The protective effect of fruits and vegetables is mostly attributed to their antioxidant compounds that neutralize natural radicals such as vitamins E and C, E (alpha-tocopherol), alpha and beta carotene, and glutathione. Pigmented fruits and vegetables that contain beta and alpha carotenoids, lutein, lycopene, zeaxanthin, astaxanthin,  $\beta$ -cryptoxanthin act as a precursor of vitamin A in the human body and in addition to compensating for vitamin deficiency in the body, play effective roles as anti-oxidants, suppressing inflammatory responses, modulating cell signaling, and inducing apoptosis. Also, they can play a protective role in cardiovascular disease, cancer, gastrointestinal disorders, and metabolic diseases [31]. Therefore, diabetic patients may need more antioxidants, and taking supplements such as vitamins D and C can have a critical role in strengthening the system of antioxidant protection and improving the lives of diabetic patients.

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**Abbreviations:** MDA: malondialdehyde, TAC: total antioxidant capacity, DM: Diabetes mellitus, BMI: body mass index, EDTA: Ethylenediaminetetraacetic acid, HPLC: high-pressure liquid chromatography, SD: Standard deviation, Vit D: Vitamin D.

**Authors' contribution:** Study concept and design: Maryam Barghi, Amir Sadeghipoor Ranjbar; Analysis and interpretation of data: Homa Moazen, and Narges Eskandari-Roozbahani; Drafting of the manuscript: Maryam Barghi, Amir Sadeghipoor Ranjbar, Narges Eskandari-Roozbahani; Critical revision of the manuscript for important intellectual content: Narges Eskandari-Roozbahani, Maryam Barghi; Statistical analysis: Homa Moazen, and Narges Eskandari-Roozbahani

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