Vitamin D Deficiency in Non-Autoimmune Hypothyroidism: A case-control study

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Research article

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Abstract

Objectives: Although in many studies, the relationship between autoimmune hypothyroidism (Hashimoto) and Vitamin D deficiency was shown, no research has been performed on the role of vitamin D in non-autoimmune hypothyroidism. Design: This was a Retrospective case–control study in Endocrinology clinic of Jahrom (south of Iran). The patients with Hashimoto (n=633) and non-Hashimoto hypothyroidism (n=305), along with a control group (n=200) were evaluated. 25(OH)D level, T3 and T4 levels were studied and Anti TPO and Anti TG tests were performed. The results of vitamin D level were analyzed and interpreted using SPSS in terms of the cause of hypothyroidism (immune and non-immune). Results: The results of the study showed a significantly lower level of vitamin D in both immune and non-immune Hashimoto's thyroiditis (HT) in comparison to healthy controls (P<0.05). We observed a significant inverse correlation between the vitamin D and TGAb level (p=0.001, r=-0.261) and a direct correlation of vitamin D with TSH level (p=0.008, r=0.108) in HT patients. Conclusion: Finally, the results indicated that non-autoimmune hypothyroidism, as well as HT, is associated with vitamin D deficiency. The role of vitamin D deficiency in HT was thought to be in the association of higher autoantibody (TGAb) level; while, there should be further studies determining vitamin D deficiency's role in non-immune hypothyroidism.

Background

Vitamin D receptors exist in many body organs. Through these receptors, vitamin D has various functions, including the regulation of ion homeostasis, cell growth, cell differentiation, and cellular immunity [1]. Vitamin D plays an important role in preventing the occurrence of many inflammatory diseases, infections, and autoimmune diseases [2]. In numerous studies, the relationship between vitamin D deficiency and a variety of diseases, including musculoskeletal [3], cardiovascular [4], kidney disease [5], diabetes [6] and infections [7] had been shown. The thyroid gland is also one of the organs that have a receptor for vitamin D. The vitamin D receptor in the thyroid is a member of a large group of receptors called nuclear receptors, which also belong to the thyroid hormones receptor [8]. Some studies indicated that vitamin D deficiency is associated with various autoimmune diseases [9]. Today, Hashimoto is one of the most common acquired hypothyroidism and autoimmune disease in children and adults [10]. The onset of autoimmune-thyroid disease with vitamin D deficiency is very common [11]. Plenty of evidence has shown the role of vitamin D in the regulation of pro-inflammatory cytokines, regulatory T cell, and immune response [12]. It seems that vitamin D deficiency leads to an increase in the risk of autoimmune diseases. Vitamin D also can reduce the pathogenesis of DCs cells, macrophage, CD4+T, CD8+T, and B cells [9]. Besides, it has been shown as a selective immune inhibitor that plays an important role in suppressing and preventing the development of autoimmune diseases such as encephalopathy, rheumatoid arthritis, systemic lupus erythematosus, diabetes type 1, and intestinal inflammatory diseases [13-15]. Recent studies have shown the role of vitamin D deficiency in autoimmune thyroid, such as HT [16, 17].

Objective:

To the best of our knowledge, there is contradictory research about the relationship between thyroid diseases, especially hypothyroidism and vitamin D deficiency; therefore, in the present study, we aimed at evaluating the vitamin D level in hypothyroidism patients. Besides, there was no study comparing the vitamin D level in
immune and non-immune hypothyroidism, and the relationship between the anti-TPO level and vitamin D, as well as the disease treatment status and vitamin D; hence, in the present study we evaluated the mentioned issues.

**Methods**

**Outcome definition:**

Normal thyroid function was considered as $0.3 \text{ mIU}/L \leq \text{TSH} \leq 3.6 \text{ mIU}/L$. The diagnosis of overt and subclinical hypothyroidism respectively was done based on TSH levels higher than 10 and $3.6 \text{ mIU}/L < \text{TSH} \leq 10 \text{ mIU}/L$ [18]. Normal T4 levels were considered between 4.5 and 12.0 μg/dL for normal participants. T4 value lower than 4.5 was one of the additional criteria' for hypothyroidism patients [31]. The values higher than 40 and 100 IU/mL were considered positive for TPOAb and TGAb, respectively. Diagnosis criteria for HT included decreased T4 value along with an elevated TSH (Overt and subclinical hypothyroidism patients) and the presence of high serum TPOAb or TGAb concentrations. The patients having overt or subclinical hypothyroidism without positive TPOAb or TGAb were considered as having non-autoimmune hypothyroidism disease. Vitamin D levels lower than 8 ng/mL were considered as severe vitamin D deficiency, 9 – 15 ng/mL concentrations as mild vitamin D deficiency, higher than 16 to 20 ng/mL concentrations as vitamin D insufficiency and higher than 20 ng/mL concentrations as normal vitamin D level [32].

**Laboratory measurements:**

Blood samples were taken from all participants after at least 8 hours of fasting. T3, Free T4, TSH were measured by Cobas ECLIA*s (Roche Diagnostics GmbH, Mannheim, Germany). Thyroid peroxidase antibody (TPOAb) were determined by chemiluminiscenta IMMULITE 2000 XPi (Siemens, Eschborn, Germany). Thyroid globulin antibody (TGAb) levels were analyzed by Enzyme-Linked Immunosorbent Assay (ELISA kit, Diesel). Vitamin D levels were measured by LIAISON vitamin D chemiluminescence immunoassay (DiaSorin, Saluggia, Italy).

**Statistical methods:**

In order to compare the quantitative continuous variables, ANOVA for parametric data and Man-u withney and Kruskal Wallis for non-parametric data were used. Chi-square test was used to compare discrete data among different groups. A p-value of less than 0.05 was considered statistically significant. SPSS v.19 was used for statistical analysis.

**Findings**

Totally 1138 individuals were studied. Demographic information and biochemical parameters of participants are presented in Table 1. Total vitamin D level of participants was 15.4(8.41-25.87). male participants had a higher level of vitamin D (p=0.001). There wasn’t any significant difference in the age of participants of Immune Hypothyroid, Non-Immune Hypothyroid and Control groups (p=0.630). Also, the distribution of male and female participants didn’t differ between there groups of the study (p=0.751).
Kruskal Wallis test revealed, there was a significant difference in Vitamin D level of study groups. As shown in figure 2., Control subjects had significantly higher vitamin D level than both HT (p=0.001) and Non-autoimmune thyroid disease patients (p=0.001), but there wasn't any significant difference between vitamin D level of HT and Non-Immune Hypothyroid patients (p>0.05).

To investigate the relationship of the thyroid autoimmunity in HT patients and vitamin D, a chi-square test investigated the distribution of the vitamin D deficiency occurrence in TPOAb+ and TGAb+ patients. The results revealed a significantly higher rate of TPOAb+ in vitamin D deficient patients (p=0.030), however, there weren't any differences in the occurrence of TGAb+ in vitamin D deficient and sufficient patients (p=0.14).

The spearman test revealed that in HT patients there was a significant inverse correlation between the vitamin D and TGAb level (p=0.001, r=-0.261) and a direct correlation of vitamin D with TSH level (p=0.008, r=0.108). However, there wasn't any significant correlation between vitamin D and other paraclinical findings). Also, these correlations were not statistically significant in Non-Immune Hypothyroid and Control.

**Discussion**

Hypothyroidism is a dysfunction in the central axis of the hypothalamus-pituitary-thyroid and insufficient secretion of the regulatory hormones is required by the thyroid gland [19]. Hypothyroidism has various reasons that the most common causes are Hashimoto or autoimmune thyroid problems and iodine disorders [20]. Previous studies have been explained pathophysiology involved in the role of vitamin D in preventing autoimmune thyroid disease. Vitamin D prevents aberrant immune responses by modulating the response of the immune cells [21]. However, Vitamin D may affect thyroid function in other ways than modulating the immune system and preventing autoimmune diseases. a limited number of the studies were performed in the field of evaluating vitamin D effect in non-immune hypothyroidism disease.

The results of the present research indicated that hypothyroidism patients with or without an immune base, deal with vitamin D deficiency (<20ng/mL) more than healthy people. In the study of Evliyaoğlu et al. [22], the patients with <20ng/mL vitamin D level was considered as vitamin D deficient and they showed that the prevalence of vitamin D deficiency is more common in people with Hashimoto than normal people. The results of their study were in agreement with those of the present study and other similar researches [16, 17, 23]. In the present study, a remarkable finding was the association between non-autoimmune hypothyroidism and vitamin D deficiency. It means that the vitamin D deficiency level in a patient with non-Hashimoto hypothyroidism was higher than the control group. The relationship between vitamin D deficiency and HT is well documented. The prevalence of positive TPOAb among people with <20ng/mL vitamin D level was more than people with sufficient vitamin D.

Krysiak et al. [24] indicated that daily uptake of 2000 IU and vitamin D can improve the treatment process in women with HT. Of course, the relationship between Hashimoto and vitamin D may also be affected by the patient’s condition. Some scholars believe that there are some differences in the past time since the onset of the disease. Effraimidis et al. [25] stated that there is no relationship between vitamin D level and primary stage of autoimmune thyroid disease. However, our subjects had crossed the initial stage of the disease. Age and sex differences may also be effective. In the present research, the male had a higher level of vitamin D than women.
Colbay et al. showed the negative correlation between TSH and vitamin D level [26]. Zhang et al. [27] indicated that the higher level of vitamin D leads to a reduction of circulated TSH, which are not in agreement with the results of our study. But in non-immune hypothyroidism vitamin D didn't correlate with TSH. This shows the diverse probable role of vitamin D deficiency in the pathogenesis of HT or non-immune hypothyroid.

The subjects of the present study were under Levothyroxine therapy, as an effective factor on TSH level, interpreting the results based TSH and finding the relationship between vitamin D and thyroid function was not reliable. Barsony et al [28] demonstrated that the treatment of hypothyroidism with Levothyroxine led to an increase in vitamin D level. While Cayir et al. showed that long-term consumption of levothyroxine may disrupt the concentration of vitamin D [29]. Therefore, there is a complicated relationship between them. Besides, similar clinical and laboratory studies did not address this issue, and they only showed the relationship between HT and vitamin D. Fournier et al. induced autoimmune thyroid disease on laboratory mice model and showed the positive effect of vitamin D on suppressing of immune system and prevention of thyroid autoimmune damage [30]. However, the present study showed that vitamin D deficiency not only affects the immune system but also have a relationship with the function of the thyroid gland.

One of the strong points of the present study is the evaluation of vitamin D level in hypothyroidism patients who their disease wasn’t autoimmune. In the present study, TGAb and TPOAb antibodies were evaluated in thyroid patients, but other factors were not evaluated. The different stage of disease in participants was the limitation of the study. For future researches, we suggest matching factors such as the stage of the disease and the dosage of daily received drugs.

**Conclusions**

The results of the present study showed that both autoimmune and non-autoimmune HT is associated with vitamin D deficiency. The relationship of vitamin D with non-autoimmune hypothyroidism is one of the new findings of the present research. In HT patients, higher vitamin D level was correlated with lower TGAb level which indicated the probable immunosuppressing role of the vitamin D.

**Results**


26. Colbay M, Altay M, Akturk M, Cakir N, Yetkin I, Arslan M. Vitamin D levels are associated with serum TSH levels but not with thyroid autoantibodies.


**References**

**Ethics approval and consent to participate:** The study was approved by the ethics committees of Jahrom University of medical sciences and written informed consent was acquired from all study subjects.

**Consent to publish:** every individual person participating in this study has signed the written informed consent of publication of their medical information freely in internet.

**Availability of data and materials:** There isn't any additional data but the data sets are available on the internal medicine department of Jahrom University of medical sciences and would be shared with anyone providing reason for using dataset by contacting the corresponding author.

**Competing interests:** None.

**Funding:** We didn't received any grant from funding agencies.
**Authors' Contributions:** The manuscript has been read and approved by all authors. All persons listed as authors have contributed to preparing the manuscript. SA designed the study and sample collecting was done by SA and MD. SA and NH undertook data analysis and writing the manuscript and reversion.

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### Tables

**Table 1- basal characteristic of patients**

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Immune Hypothyroid</th>
<th>Non-Immune Hypothyroid</th>
<th>control</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number</td>
<td>633</td>
<td>305</td>
<td>200</td>
<td></td>
</tr>
<tr>
<td>Age (IQR)</td>
<td>37.48±13.18</td>
<td>36.65±14.56</td>
<td>37.69±15.26</td>
<td>0.630</td>
</tr>
<tr>
<td>SEX (male), n (%)</td>
<td>146 (23.1)</td>
<td>66 (21.6)</td>
<td>49 (24.5)</td>
<td>0.751</td>
</tr>
<tr>
<td>Vitamin D level, ng/ml (IQR)</td>
<td><strong>13.22(8.1-24.27)</strong></td>
<td><strong>16(8.43-28.85)</strong></td>
<td><strong>20.4(11.2-29.6)</strong></td>
<td>0.001</td>
</tr>
<tr>
<td>TSH, mIU/l (IQR)</td>
<td>6.29(3.13-17.75)</td>
<td>5.92(2.54-1.81)</td>
<td>2.2(1.59-2.88)</td>
<td>0.001</td>
</tr>
<tr>
<td>T3, mg/ml (IQR)</td>
<td>1.67(1.26-2.38)</td>
<td>1.68(1.22-2.15)</td>
<td>1.32(0.74-1.8)</td>
<td>0.001</td>
</tr>
<tr>
<td>T4, mg/ml (IQR)</td>
<td>9.1(7.06-63.62)</td>
<td>11.1(7.42-80.05)</td>
<td>7.05(0.63-808)</td>
<td>0.001</td>
</tr>
<tr>
<td>TPOAb, IU/mL (IQR)</td>
<td><strong>14.4(4.69-134.6)</strong></td>
<td>3.59(1.13-16.67)</td>
<td><strong>19.4(13.6-33.55)</strong></td>
<td>0.001</td>
</tr>
<tr>
<td>TGAb, IU/mL (IQR)</td>
<td>320.7(112.2-733)</td>
<td>10(4.33-15.63)</td>
<td><strong>16.9(9.16-185.95)</strong></td>
<td>0.001</td>
</tr>
</tbody>
</table>

**Table 2. Thyroid autoimmunity in HT patients**

<table>
<thead>
<tr>
<th>Vitamin D Sufficient (&gt;20 ng/ml)</th>
<th>TPOAb+</th>
<th>TGAb+</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Deficient (&lt;20 ng/ml)</td>
<td>422</td>
<td>34</td>
<td>0.030</td>
</tr>
</tbody>
</table>

### Figures
Figure 1

Study participants
Figure 2

Vitamin D level of study groups