Vitamin B12 and Vitamin D Levels in Patients with Autoimmune Hypothyroidism and Their Correlation with Anti-Thyroid Peroxidase Antibodies

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Significance of the Study

- There was a negative correlation between vitamin B12 and vitamin D levels and anti-thyroid peroxidase antibodies in patients with autoimmune hypothyroidism.

Keywords

Vitamin B12 · Vitamin D · Autoimmune hypothyroidism · Anti-thyroid peroxidase antibodies

Abstract

Background: This study was designed to investigate the vitamin D (vit-D) and vitamin B12 (vit-B12) levels and their correlation with anti-thyroid peroxidase (anti-TPO) antibodies in patients with autoimmune hypothyroidism. **Methods:** A total of 130 patients diagnosed with autoimmune hypothyroidism were included in the study retrospectively. The patients were divided into two groups as those having vit-B12 levels below 200 pg/mL ($n = 60$) and vit-B12 levels equal to or above 200 pg/mL ($n = 70$). These two groups were compared in terms of age, sex, thyroid-stimulating hormone (TSH), free-T4 (FT4), vit-D, and anti-TPO. The correlation between vit-B12 and anti-TPO levels was also investigated in these groups. Patients were then divided into four groups according to their vit-D levels. Patients with normal vit-D levels ($25[\text{OH}]\text{D} > 30 \text{ ng/mL}$; $n = 5$), those with vit-D insufficiency ($20\text{--}30 \text{ ng/mL}$; $n = 9$), those with vit-D deficiency ($10\text{--}20 \text{ ng/}$

mL ; $n = 43$), and those with severe vit-D deficiency ($< 10 \text{ ng/mL}$; $n = 73$). These four groups were compared in terms of age, gender, TSH, FT4, vit-B12, and anti-TPO levels. In addition, the correlation between levels of vit-D and anti-TPO was also investigated. **Results:** We found that vit-B12 deficiency and vit-D deficiency were associated with autoimmune hypothyroidism, and that there was a negative correlation between vit-B12 and vit-D levels and anti-TPO antibodies in these patients. **Conclusion:** In patients with autoimmune hypothyroidism, vit-D and vit-B12 deficiency should be investigated at the time of diagnosis and periodically on follow-ups.

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Published by S. Karger AG, Basel

Introduction

Hashimoto thyroiditis (HT) is the most common cause of hypothyroidism, a disease caused by inadequate synthesis and/or release of thyroid hormones [1]. In these patients, hypothyroidism at various levels is accompanied by autoantibodies (antithyroid peroxidase antibody

[anti-TPO], antithyroglobulin antibody) and lymphocytic infiltration in thyroid tissue. It is more common in the 30- to 50-year-old age group and is 4–10 times more common in women than in men [2].

Environmental (30%) and genetic (70%) factors influence the etiopathogenesis of HT [3]. Genetic factors include the major histocompatibility gene (*HLA*), immune regulator genes (*CTLA4*, *PTPN22*, *FOXP3*, *CD25*, *CD40*, and *FRCL3*) and thyroid-specific genes (*TSHR*, *Tg*) [3]. The possible environmental factors in etiology are smoking, alcohol, dietary iodine, stress, selenium, vitamin D (vit-D) deficiency, bacterial and viral infections, pregnancy, and drugs [4].

The main function of vit-D is to achieve calcium and phosphorus balance with bone metabolism. In recent years, however, vit-D has been reported to have a role in the formation of autoimmune diseases, heart diseases, cancer, inflammatory bowel diseases, diabetes, and rheumatologic diseases, as well as its effects on bone metabolism [5]. In addition, it is thought that vit-D may play a role in the regulation of the immune system because vit-D receptors are detected on cells of the immune system [6]. In some studies, a relationship between vit-D deficiency and autoimmune thyroid diseases such as HT and Grave's disease has been reported, and impaired vit-D signaling has been reported in thyroid cancers [7]. Bozkurt et al. [8] reported that vit-D deficiency had a role in the development of HT.

Many autoimmune diseases, such as pernicious anemia, myasthenia gravis, idiopathic hypoparathyroidism, vitiligo, celiac disease, type 1 diabetes mellitus, autoimmune liver diseases, primary biliary cirrhosis, multiple sclerosis, Addison's disease, rheumatoid arthritis, and systemic lupus erythematosus, may accompany HT [9]. Vitamin B12 (vit-B12) deficiency has been reported frequently in autoimmune thyroid patients [10]. This association is probably due to impaired absorption of vit-B12 by atrophic gastritis and/or pernicious anemia associated with autoimmune thyroid disease [11]. Atrophic gastritis is seen in 35–40% of autoimmune thyroid diseases [12]. It was reported that 1/3 of the patients with primary hypothyroidism had antibodies against gastric parietal cells and 12% had pernicious anemia. Intrinsic factor antibodies have also been shown in these patients [13].

HT is an autoimmune disease and studies suggest that vit-D deficiency has an effect on the etiopathogenesis of the disease. In addition, vit-B12 deficiency is frequently seen in this disease [7, 10]. This study was designed to investigate levels of vit-D and vit-B12 in patients with autoimmune hypothyroidism and to evaluate their correlation with anti-TPO antibodies.

Table 1. Demographic data and distribution of parameters

Male	15 (11.5)
Female	115 (88.5)
Vit-B12 <200 pg/mL	60 (46)
Vit-D <30 ng/mL	125 (96)
Anti-TPO (+)	130 (100)
Age, years	
Mean	41.4±1.9
Median	41 (18–64)

Values are presented as *n* (%), the mean ± SD, or median (min.–max.).

Materials and Methods

A total of 130 patients aged between 18 and 65 years who were diagnosed with autoimmune hypothyroidism in the Internal Medicine Outpatient Clinic of Umraniye Education and Research Hospital were included in the study retrospectively. Approval was obtained from the Ethics Committee of our hospital. Exclusion criteria were thyroidectomy, receiving RAI treatment, malabsorption syndromes, gastrectomy, ileal resection, a vegetarian diet, pregnancy, and use of drugs to treat vit-B12 deficiency (phenytoin, metformin, and DHF reductase inhibition).

The patients were divided into two groups as those having vit-B12 levels below 200 pg/mL (*n* = 60) and vit-B12 levels equal to or above 200 pg/mL (*n* = 70) [14]. These two groups were compared in terms of age, sex, thyroid-stimulating hormone (TSH), free-T4 (FT4), vit-D, and anti-TPO. In addition, the correlation between levels of vit-B12 and anti-TPO was also examined in these two groups. Patients were then divided into four groups according to their vit-D levels [5]: patients with normal vit-D levels (25[OH]D >30 ng/mL; *n* = 5), those with vit-D insufficiency (20–30 ng/mL; *n* = 9), those with vit-D deficiency (10–20 ng/mL; *n* = 43), and those with severe vit-D deficiency (<10 ng/mL; *n* = 73). Then the age, gender, TSH, FT4, vit-B12, and anti-TPO levels of these four groups were compared. In addition, the correlation between vit-D and anti-TPO levels was also investigated in these four groups.

Serum vit-B12 (187–883 pg/mL), vit-D (>30 ng/mL), TSH (0.35–4.94 mU/mL), FT4 (0.4–1.48 ng/mL), and anti-TPO (<5.6 IU/mL) measurements were performed in the fasting blood taken from the patients on the same day. These tests were performed by ABBOTT Architect i2000 with the chemiluminescence method. The anti-TPO antibody cut off level was accepted as 5.61 IU/mL.

Statistical Analysis

Statistical analyses were performed using the MedCalc Statistical Software version 12.7.7 (MedCalc Software bvba, Ostend, Belgium; <http://www.medcalc.org>; 2013). Statistical significance was set at *p* < 0.05. Descriptive statistics were used to define continuous variables (mean ± standard deviation, and median, minimum and maximum). The comparison of two independent groups with a normal distribution was performed using the Student *t* test. The comparison of two independent groups without a normal distribution was performed using the Mann-Whitney U test. The correlation

Table 2. Relationship between vit-B12 levels and TSH, FT4, anti-TPO, and vit-D

	Vit-B12 <200 pg/mL (<i>n</i> = 60)		Vit-B12 ≥200 pg/mL (<i>n</i> = 70)		<i>p</i>
TSH, mU/mL	14.9±18.5	9.5 (4–100)	13.4±10.3	8.0 (5–45)	0.302 ^a
FT4, ng/mL	1.0±1.3	0.8 (0.2–8.0)	1.0±1.0	0.9 (5–6.5)	0.614 ^a
Anti-TPO, IU/mL	484.7±377.1	460 (4–1000)	292±389.4	34 (5–1000)	<0.001 ^a
Vit-D, ng/mL	11.9±7	10 (4.1–36)	11.5±7.7	9 (3.6–36)	0.186 ^a
Age, years	41.4±12	39 (19–64)	41.3±11.9	41 (18–64)	0.967 ^b
Male	8 (13.3)		7 (10)		
Female	52 (86.7)		63 (90)		

Data are presented as the mean ± SD, median (min.–max.), and *n* (%).

^a Mann-Whitney U Test. ^b Student *t* test.

Fig. 1. Correlation between anti-TPO and vit-B12. There was a weak, negative, statistically significant correlation between anti-TPO and vit-B12 levels in both groups with and without vit-B12 deficiency ($r = -0.394$, $p < 0.001$).

tion between two continuous variables not compatible with a normal distribution was examined with the Spearman rho correlation coefficient. Statistical evaluation of categorical variables was performed using the χ^2 test.

Results

The mean age of all patients was 41.4 ± 11.9 years, and 115 patients (88.5%) were female and 15 (11.5%) were male. Vit-B12 deficiency was found in 60 (46%) and vit-D deficiency was found in 125 (96.1%) of all patients. Seventy-three patients (56.1%) with hypothyroidism had severe vit-D deficiency. Anti-TPO antibody was positive in all patients (Table 1).

There was no significant difference between the two vit-B12 groups in terms of age, gender, TSH, FT4, and vit-D levels. However, the anti-TPO levels were significantly higher in patients with low vit-B12 levels ($p < 0.001$; Table 2). There was no significant difference between the four vit-D groups in terms of age, gender TSH, FT4, vit-B12, and anti-TPO levels (Table 3).

There was a weak, negative correlation between TSH and vit-D levels in the $25(\text{OH})\text{D} < 10$ ng/mL group. Moreover, there was also a weak, negative correlation between anti-TPO and vit-D levels in the $25(\text{OH})\text{D} < 10$ ng/mL group. There was a weak, negative correlation between anti-TPO and vit-B12 levels in all patient groups ($p < 0.001$; Table 4; Fig. 1). There was also a weak, negative correlation between anti-TPO and vit-D levels in all of the groups (Table 5; Fig. 2).

Discussion

HT is the most common cause of primary hypothyroidism. The disease is more common in the 30- to 50-year age group, and 95% of patients are women. The female to male ratio is about 7.2 [15]. In our study, the male to female ratio was 7.7, and the mean age of the patients was 41.4 ± 11.9 years, similar to the literature.

There are several studies suggesting that the prevalence of vit-D deficiency is high in patients with HT and that there is a relationship between hypothyroidism and vit-D in these patients [16]. In our study, we found that 96% of patients with HT had vit-D deficiency, and 56.1% of patients had severe vit-D deficiency (<10 ng/mL). We also found a negative correlation between vit-D levels and anti-TPO levels in these patients. We think that vit-D deficiency may cause autoimmune hypothyroidism. Bozkurt et al. [8] reported that vit-D deficiency had a role in the development of HT.

Some studies have reported a relationship between vit-D deficiency and thyroid autoimmunity in all age groups, and decreased anti-TPO levels when cholecalciferol supplementation is performed in HT patients with vit-D deficiency [17, 18]. Kivity et al. [19] reported that the prevalence of vit-D deficiency (25[OH]D level <25 nmol/L) was higher in people with autoimmune thyroid disease. They also reported a correlation between vit-D deficiency and anti-TPO levels and that this had an effect on the pathogenesis of autoimmune thyroid diseases. Mansournia et al. [20] reported a significant negative correlation between HT disease and serum levels of 25(OH)D. Shin et al. [21] reported that people with high anti-TPO levels had lower vit-D levels.

Mazokopakis et al. [17] showed that serum levels of 25(OH)D were inversely related to anti-TPO levels in patients with HT. Anti-TPO levels were also significantly higher in HT patients with vit-D deficiency (<75 nmol/L) than those HT patients without vit-D deficiency. In another study, a significant decrease in serum anti-TPO levels was reported after 4 months of oral vit-D3 supplementation (1,200–4,000 IU/day) in patients with vit-D deficiency [17]. Screening for vit-D deficiency and replacement therapy if necessary has been recommended [22]. Levels of vit-D were observed to be significantly lower in children with HT [23]. A negative correlation between vit-D and TPO antibodies was found in our study. HT disease activity has been reported to decrease with vit-D treatment [24].

The relationship between vit-D deficiency and thyroid autoimmunity remains unclear. However, it is suggested

Table 3. The relationship of vit-D levels with parameters

	Vit-D <10 ng/mL (n = 73)	Vit-D 10–20 ng/mL (n = 43)	Vit-D 20–30 ng/mL (n = 9)	Vit-D >30 ng/mL (n = 5)	p ^a
TSH, mU/mL	15.2±17.3	12.9±10.8	12.4±9.1	11.6±10.4	0.877
FT4, ng/mL	1.0±1.0	1.1±1.5	0.9±0.1	0.8±0.1	0.579
Anti-TPO, IU/mL	411.9±413.3	284.9±323.1	450.6±505.1	487.8±472.4	0.429
Vit-B12, pg/mL	220.6±70.1	208.6±59.1	210±42.6	238.2±87.6	0.753
Age, years	41.0±11.8	42.2±12.9	42.6±11.1	37.4±3.4	0.812
Vit-D <10 ng/mL					
Vit-D ≥10 ng/mL					
Gender and vit-D relationship					
Male	5 (6.8)				
Female	68 (93.2)	10 (17.5)			
Total	73 (100)	47 (82.5)			
		57 (100)			

Data are presented as the mean ± SD, median (min.–max.), and n (%).
^a Kruskal-Wallis test.

Table 4. Correlation between levels of vit-B12 and other parameters

	r^a	p
Vit-B12 <200 pg/mL ($n = 60$)		
TSH and B12	-0.182	0.164
Anti-TPO and B12	-0.170	0.194
FT4 and B12	-0.055	0.674
Vit-B12 $\geq$ 200 pg/mL ($n = 70$)		
TSH and B12	-0.102	0.401
Anti-TPO and B12	-0.170	0.159
FT4 and B12	0.062	0.609
All groups		
TSH and B12	0.018	0.835
Anti-TPO and B12	-0.394	<0.001
FT4 and B12	0.039	0.658

^a Spearman rho correlation coefficient.

Table 5. Correlation between vit-D and other parameters

	r^a	p
Vit-D <10 ng/mL ($n = 73$)		
TSH and vit-D	-0.304	0.009
Anti-TPO and vit-D	-0.257	0.028
FT4 and vit-D	0.177	0.134
Vit-D $\geq$ 10 ng/mL ($n = 57$)		
TSH and vit-D	-0.021	0.876
Anti-TPO and vit-D	0.067	0.619
FT4 and vit-D	0.005	0.969
All groups		
TSH and vit-D	-0.149	0.091
Anti-TPO and vit-D	-0.185	0.035
FT4 and vit-D	0.068	0.441

^a Spearman rho correlation coefficient.

Fig. 2. Correlation between anti-TPO and vit-D. There was also a weak, negative, and statistically significant correlation between anti-TPO and vit-D levels in all these groups ($r = -0.185$, $p = 0.035$).

that vit-D may be associated with anti-inflammatory and immunomodulatory effects. Vit-D plays a significant role in modulation of the immune system, enhancing the innate immune response while exerting an inhibitory action on the adaptive immune system. Most immune cells, including T cells, B cells, dendritic cells, and macrophages, express the vit-D receptor and 1α -hydroxylase [25].

In our study, we found that the incidence of vit-B12 deficiency in autoimmune hypothyroid patients was 46%. We also found that there was a negative correlation between anti-TPO and vit-B12 levels. It is possible that this may be due to pernicious anemia or atrophic gastritis accompanying autoimmune hypothyroidism, which is often accompanied by other autoimmune diseases. Perni-

cious anemia may be part of a polyglandular endocrinopathy associated with hypothyroidism [26]. This association is caused by atrophic gastritis and/or pernicious anemia associated with autoimmune thyroid diseases disrupting vit-B12 absorption. Overall, 35–40% of patients with autoimmune thyroid diseases have atrophic gastritis [26], while 33% of patients with primary hypothyroidism have been shown to have antibodies to gastric parietal cells, and 12% have pernicious anemia. Intrinsic factor antibodies have also been observed in patients with autoimmune hypothyroidism [9].

The prevalence of vit-B12 deficiency in patients with hypothyroidism in Pakistan, Turkey, and India is reported to be 40.5, 18.6, and 10%, respectively [27]. The differences in prevalence between these ratios could be caused

by differences in diet and nutrition status. Clinical manifestations of vit-B12 deficiency are very variable. Hematological and neurological abnormalities may coexist, be independently found, or be very uncertain. Even in asymptomatic patients, the first sign may be the detection of vit-B12 deficiency [28]. Patients with both hypothyroidism and vit-B12 deficiency also have similar symptoms. Symptoms such as fatigue, weakness, dementia, depression, memory impairment, lethargy, and tingling are frequently seen in patients with hypothyroidism and vit-B12 deficiency. Thus, vit-B12 deficiency may be ignored in hypothyroidism.

Vit-B12 deficiency increases homocysteine levels [13] and a relationship is known to exist between hyperhomocysteinemia and the development of atherosclerosis. Several studies have shown a relationship between hypothyroidism and homocysteine levels. Thus, if vit-B12 deficiency is overlooked in hypothyroid patients it may increase cardiovascular risk through hyperhomocysteinemia [29].

Ness-Abramof et al. [30] recommend screening vit-B12 levels at the time of diagnosis of autoimmune thyroid disease, with 3- to 5-year intervals. Collins and Pawlak [28] also recommend screening for vit-B12 deficiency in hypothyroid patients at the time of diagnosis and at certain intervals periodically. Our study supports these recommendations.

One of the limitations of this study is its retrospective design. Another is the small number of patients included. Studies investigating the relationship between future hypothyroidism and vit-D and vit-B12 should be prospectively designed in larger sample sizes.

Conclusion

In this study, we observed vit-B12 and vit-D deficiency in patients with HT. We also found a negative correlation between vit-D deficiency, vit-B12 deficiency, and levels of anti-TPO. We suggest that vit-D deficiency contributes to the etiopathogenesis of HT, and that vit-B deficiency and HT are part of a polyglandular endocrine pathology. Vit-D and vit-B12 deficiency should be investigated in patients with autoimmune hypothyroidism at the time of diagnosis and during periodic follow-ups, and supplementation should be performed in those who need it. More randomized controlled trials are needed to investigate the effect of vit-D and vit-B12 deficiency in the etiopathogenesis of HT, and to determine the effect of replacement therapy on the clinical course of HT.

Statement of Ethics

All procedures performed in this study involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Disclosure Statement

All authors declare that they have no conflicts of interest. No financial support was obtained.

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