Case Report

Hashimoto’s disease, which is associated with thyroid ophthalmopathy: a case report

Tiroid oftalmopatisi ile ilişkili Hashimoto hastalığı

Cem Onur Kirac 1*, Suleyman Hilmi Ipekci 1, Suleyman Baldane 1, Banu Turgut Ozturk 2, Levent Kebapcilar 1

1 Selcuk University, Faculty of Medicine, Department of Internal Medicine, Division of Endocrinology and Metabolism, Konya, Turkey
1 Selcuk University, Faculty of Medicine, Department of Ophthalmology, Konya, Turkey
* Corresponding author: Cem Onur Kirac E-mail: cokirac@gmail.com ORCID: 0000-0002-0249-9867
Received: 17 September 2019 Accepted: 22 November 2019

ABSTRACT

Hashimoto’s disease is the most common cause of hypothyroidism. Since it is an autoimmune disease, the incidence of other autoimmune diseases has increased with Hashimoto’s disease. Extra-thyroidal involvements are also seen due to the autoantibodies that play a role in the etiopathogenesis of the disease. One of these is ophthalmopathy. We aimed to report the association of ophthalmopathy with Hashimoto’s disease which is frequently associated with Graves’ disease and to report the mechanism involved in pathogenesis in the light of literature.

Keywords: Etiopathogenesis, Hashimoto’s disease, thyroid ophthalmopathy, thyroid stimulating immunoglobulin

ÖZ


Anahtar kelimeler: Etiopathogenez, Hashimoto hastalığı, tiroid oftalmopatisi, tiroid stimulan antikor
INTRODUCTION

Thyroid-associated ophthalmopathy (TAO) is an autoimmune disease characterized by thickening of extraocular muscles, increased fat and connective tissue [1]. Although the pathogenesis of TAO has not been completely understood yet, it is the hypothesis that the proteins produced by orbital fibroblasts are presented as autoantigen and the autoimmune response to these autoantigens may take part in a major role of the disease’s pathogenesis [2]. Some studies have shown that ectopic thyrotropin receptors expressed by orbital fibroblasts and adipocytes contribute to this inflammatory process [3]. While TAO is frequently associated with hyperthyroidism (90%), 6% of these patients are euthyroid and 1% may be hypothyroid. Three percent of patients are diagnosed as Hashimoto’s thyroiditis (HT) at the time of diagnosis with TAO [3]. In this case, we aimed to keep HT in mind in patients with suspected TAO.

CASE

A 37-year-old female patient who was diagnosed with hypothyroidism was admitted to the ophthalmology clinic with the complaint of a lower-than-right on her left eyelid, which she noticed for the last few weeks. In the patient’s examination, bilateral visual acuity, anterior segment and fundus findings were normal. The left eyelid was in the normal position, but the right eyelid was 2 mm above the limbus, and the right eyelid retraction was detected (Figure 1). Hertel exophthalmometer measurement was 19 mm on the right and 20 mm on the left (BC 102). Orbital MRI showed a thickening of the right rectal rectus muscle (Figure 2). The patient was referred to the Endocrinology clinic with the suspicion of thyroid ophthalmopathy. Physical examination did not reveal any abnormality except for exophthalmos at her right eye. In her anamnesis, it is learned that she had received levothyroxine replacement therapy at 1.7 mcg/kg/day, for more than 10 years without any hyperthyroidism phase of her illness. There was no additional disease or other medication. Laboratory evaluation revealed that, TSH: 1.16 mU/ml (normal: 0.27-4.2); anti-thyroid peroxidase (TPO): 242 IU/ml (normal: 0-34), anti-thyroglobulin (Tg): 109 IU/ml (normal: 0-115). TSH receptor stimulant antibody (TSAb) level was measured as 4 IU/l (normal <0.10). The patient was planned to receive methylprednisolone at a dose of 250 mg/week for 4 weeks. At the end of the treatment the patient’s complaint was regressed. Written informed consent was taken from the patient.

DISCUSSION

Although HT was first described by Hakaru Hashimoto in 1912, it is still an autoimmune disease whose etiopathogenesis is not fully explained after nearly 100 years. HT is the most common autoimmune disease and endocrine disorder and also the most common cause of hypothyroidism [4-6]. There is a correlation between anti-TPO titers, lymphocyte cell infiltration and the degree of hypoechochogenicity in thyroid ultrasound [7]. In addition to anti-TPO, anti-Tg antibody, TSH receptor blocking antibody and rarely TSAb can be seen in HT [8].

The publications related to HT-associated orbitopathy are generally case-based and there are rare original studies. George et al. [9] reported that 6% of patients had TAO in their study on 700 HT diagnosed patients. While TSAb
positivity was 68.2% in patients with TAO in his study, this rate was found to be 5.5% in HT patients without ocular involvement. TSAb titer was also significantly higher in patients with TAO. Age and smoking were also related to TAO. As a result of this study, it is concluded that TSAb is responsible for HT-associated orbitopathy, as in the case of Graves’ orbitopathy. In another retrospective study, HT-related ophthalmopathy was found to be less severe than Graves-related ophthalmopathy, and this was thought to be associated with TSAb titer [10].

The most common involvement of Graves’ disease is orbitopathy [11]. Thyroid dermopathy can be seen in 4% of the patients and there is often eye involvement in these patients [12]. These two involvements are thought to have autoimmune origin and are related to TSAb. In another case, both ocular involvement and dermopathy were observed in a patient with positive TSAb with HT [13]. This case suggests that extra-thyroidal involvement may not be specific to Graves’ disease, but is directly related to TSAb positivity.

There are also cases of TSAb negative HT-related ophthalmopathy [14]. In this instance, apart from TSAb, other factors should be considered in the etiopathogenesis. However, patients’ benefit from steroid treatment as in our case still indicates that underlying inflammatory events are at the forefront.

CONCLUSION

As a result, although serum TSAb level is thought to be associated with eye involvement in HT as in Graves ophthalmopathy, other factors that may lead to the same situation should be clarified with more extensive molecular and genetic studies. Nevertheless, it should be kept in mind that patients with ophthalmopathy should evaluate for TSAb levels even if they are euthyroid, even hypothyroid.

DECLARATION OF CONFLICT OF INTEREST

The authors received no financial support for the research and/or authorship of this article. There is no conflict of interest.

REFERENCES