Transformation of hashimoto thyroiditis to Graves’ disease

Stella Pak, Damian Valencia and Adam Fershko*
Department of Medicine, Kettering Medical Center, Kettering, 3535 Southern Blvd, Kettering, OH 45429, USA

Abstract
Hashimoto’s thyroiditis and Graves’ disease are thyroid-specific autoimmune disorders with distinct pathological mechanism. Clinical presentation and histopathological features also vary greatly in these disorders. Sequential conversion from Hashimoto’s thyroiditis (HT) to Graves’ disease (GD) has been rarely reported throughout the world. Herein, we report a woman with history of HT, who then developed GD requiring total thyroidectomy. By reporting this case, we hope to raise awareness about this phenomenon among clinicians and illustrate the potential mechanisms behind such change. Improving recognition and identification of this potential conversation between HT and GD would allow clinicians to expeditiously implement appropriate treatment, increasing the quality of care.

Introduction
Hashimoto thyroiditis (HT) is a chronic inflammatory disorder of thyroid glands. HT causes hypothyroidism and affects approximately 0.8% of the population. Anti-thyroperoxidase antibodies and antithyroglobulin antibodies are found in 95% and 60-80% of patients with HT, respectively [1]. These autoantibodies cause progressive destruction of thyroid follicles, eventually resulting in atrophy and fibrosis of thyroid glands. CD4 (+) T-Helper-1 (TH-1) cells play a major role in the pathogenesis of HT [2].

Contrarily, Graves’ disease (GD), an autoimmune process which leads to hyperthyroidism, is characterized by the presence of antithyrotropin receptor antibodies and clinically by exophthalmos and pretibial myxedema. GD affects about 0.5% of the population and can possibly cause thyrotoxicosis [3,4]. In GD Autoantibodies overstimulates thyroid-stimulating hormone (TSH) receptors, resulting in hyperthyrotope of thyroid glands. Unlike HT, TH-2 cells play a key role in disease process of GD [2]. Because different autoantibodies and TH cells are involved, the conversion between HT and GD is rare; but, yet a sequential conversion of HT to GD is even more rare [4]. Herein, we report a woman with history of HT, who then developed GD that required a total thyroidectomy.

Case presentation
The patient is a 41-year-old female with a past medical history of Hashimoto thyroiditis who presented with new-onset palpitation, unintentional weight loss (approximately 10 lbs.), hair loss, fatigue, constipation and tremors over the past 5 months. On presentation, the patient vitals were: blood pressure 148/82 mmHg, heart rate 72 / min, respiratory rate 16 /min, temperature 37 °C and oxygen saturation of 98%. Initial laboratory results revealed TSH 0.02 ng/dL, T3 of 1.71 ng/dL, and T4 3.2 ng/dL, which were significantly different from her baseline (TSH 3.73 ng/dL, T3 1.08 ng/dL, and T4 2.70 ng/dL). After initial evaluation, the patient’s home levothyroxine (112 mg daily) was discontinued and she was started on propranolol (10 mg daily). From initial onset of hypothyroidism, she continued to be sero-negative for thyroid stimulating antibodies, at 3.93 IU/L, diagnostic of Graves’ disease. Her thyroid stimulating immunoglobulin level was also 164 units.

The ultrasonogram of her thyroid revealed a bilateral multinodular goiter and heterogeneous echotexture (Figure 1). On her radioactive studies, she had heterogeneous uptake. Radioiodine ablation was not possible due to her iodine allergy and the heterogeneous uptake within the gland. Thus, our patient underwent a total thyroidectomy. Pathological examination of removed thyroids revealed abundant thyroglobulin, consistent with the diagnosis of hyperthyroidism (Figure 2). She had uneventful postoperative recovery, but developed postoperative hypothyroidism. Her laboratory results were TSH 2.73 ng/dL, T3 of 0.74 ng/dL, and T4 2.70 ng/dL. As a result, the patient was started on levothyroxine (112 mg daily) and now remains symptom free.

Discussion
When the patient with HT presents with symptoms of hyperthyroidism, the most likely cause is over-replacement of thyroid hormone. However, as highlighted in the present case, it is crucial to include GD in the differential diagnosis, given the common autoimmune origin of both disorders. As GD can cause irreversible complications such as exophthalmos, timely diagnosis and treatment is important to improve and maintain the quality of lives of patients [3].

The pathophysiology underlying the conversion of HT to GD remains largely unknown. There are three plausible hypotheses for this rare transformation. First hypothesis postulates that the development of various autoantibodies can give rise to intricate interaction between multiple types of antibodies and thyroid glands, leading to the conversion. Second hypothesis is that the thyroid dysfunction presented as hypothyroidism from severe autoimmune-induced tissue injury, but then later thyroid glands gradually recover from the injury and become able to response to thyroid stimulating antibodies, leading to hyperthyroidism [5]. Third possible hypothesis is that age-related

*Correspondence to: Stella Pak, MD, Department of Medicine, Kettering Medical Center, Kettering, 3535 Southern Blvd, Kettering, OH 45429, USA, Tel. (937) 395-8997; Fax: (937) 395-8399; E-mail: Stella.Pak@ketteringhealth.org

Key words: hashimoto thyroiditis, graves’ disease, thyroidectomy

Received: October 08, 2017; Accepted: October 23, 2017; Published: October 26, 2017
Hypothyroidism from HT was traditionally believed to result from the gradual destruction of thyroid follicles by autoantibodies [7]. Thus, the sequential conversion from HT to GD challenges the current view in pathophysiology of HT. In the patients who had a shift from HT to GD, the autoantibodies could have merely blocking the TSH receptors rather than infiltrating and destructing the thyroid glands. Accumulation and analysis of cases of the rare transformation would enable unraveling of its pathophysiology. The reason for lower incidence of conversion from HT to GD than the other way around is not known [4]. The symptoms of HT tend to be mild and, thus, more likely to be overlooked than that of GD. Based on this, the transformation from HT to GD may not be as rare as inferred by literature, but, rather, missed.

In summary, we described a case of a sequential transformation from HT to GD. By reporting this case, we hope to raise the awareness about this rare phenomenon among clinicians and illustrate the potential mechanisms behind such change. Improvement of recognition and identification of this conversation process between HT and GD would allow clinicians to expeditiously implement appropriate treatment, thus increasing the quality of care.

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