Thyrotoxicosis: An unusual presentation

Somnath Gooptu, Gurjit Singh, Iqbal Ali, Siddharth Mishra

ABSTRACT

Introduction: In a thyrotoxic patient, Grave’s disease, solitary toxic nodule and toxic nodular goitre are considered to be the possible diagnosis. However, in certain inflammatory conditions also present with similar features wherein thyroid hormones are released due to destruction of the gland. Such conditions as Hashimoto’s thyroiditis may be missed on cytological examination and are diagnosed only on histopathological examination. The combination of Hashimoto’s thyroiditis with toxicity is called hashitoxicosis.

Case Report: A 37-year-old female presented with thyroid swelling and features of thyrotoxicosis which were confirmed by thyroid function tests. A clinical diagnosis of toxic nodular goitre was made. Euthyroid state was achieved after treatment with tab carbimazole and beta blockers. The patient underwent near total thyroidectomy. After surgery, patient developed hypocalcemia which was managed by intravenous and oral calcium supplementation. At postoperative estimation thyroid auto antibodies were found elevated and the patient was started on tab thyroxine. At sixth month follow-up serum calcium levels returned to normal levels and hence calcium supplementation was stopped.

Conclusion: Any patient presenting in thyrotoxic state, possibility of hashitoxicosis should be considered and confirmed by the estimation of thyroid antibodies and the use of ultrasonography guided fine-needle aspiration cytology.
CASE REPORT

Thyrotoxicosis: An unusual presentation

Somnath Gooptu, Gurjit Singh, Iqbal Ali, Siddharth Mishra

ABSTRACT

Introduction: In a thyrotoxic patient, Grave’s disease, solitary toxic nodule and toxic nodular goitre are considered to be the possible diagnosis. However, in certain inflammatory conditions also present with similar features wherein thyroid hormones are released due to destruction of the gland. Such conditions as Hashimoto’s thyroiditis may be missed on cytological examination and are diagnosed only on histopathological examination. The combination of Hashimoto’s thyroiditis with toxicity is called hashitoxicosis. Case Report: A 37-year-old female presented with thyroid swelling and features of thyrotoxicosis which were confirmed by thyroid function tests. A clinical diagnosis of toxic nodular goitre was made. Euthyroid state was achieved after treatment with tab carbimazole and beta blockers. The patient underwent near total thyroidectomy. After surgery, patient developed hypocalcemia which was managed by intravenous and oral calcium supplementation. At postoperative estimation thyroid auto antibodies were found elevated and the patient was started on tab thyroxine. At sixth month follow-up serum calcium levels returned to normal levels and hence calcium supplementation was stopped. Conclusion: Any patient presenting in thyrotoxic state, possibility of hashitoxicosis should be considered and confirmed by the estimation of thyroid antibodies and the use of ultrasonography guided fine-needle aspiration cytology.

Keywords: Grave’s disease, Hashitoxicosis, Thyroid antibodies, Thyrotoxicosis

INTRODUCTION

In a patient of thyrotoxicosis possible etiologies considered are Grave’s disease, toxic multi-nodular goitre and solitary toxic nodule. However, certain inflammatory conditions may lead to destruction and damage to thyroid gland resulting in leakage of hormones causing transient thyrotoxicosis. Such transient thyrotoxicosis in association with Hashimoto’s thyroiditis is termed hashitoxicosis. Patients with this disorder are expected to remit or even develop hypothyroidism. Only correlation of histopathology with antibody testing may clinch the diagnosis.

CASE REPORT

A 37-year-old female presented with complaints of thyroid swelling for the last three months which was associated with tachycardia and positive Stellwag’s sign with no previous history of thyrotoxicosis. Thyroid
function tests revealed normal levels of T₃ (2 nmol/L) and T₄ (14.8 nmol/L) but decreased levels of TSH (0.011 mU/L) hence diagnosis of thyrotoxicosis was made. Ultrasound of the neck revealed multinodular goitre. Fine-needle aspiration cytology (FNAC) was suggestive of colloid goitre (Figure 1).

The patient was started on antithyroid drugs and beta blockers. Euthyroid state was achieved in four weeks. Patient underwent near total thyroidectomy. Within 24 hours following surgery patient developed features of hypocalcemia. She had multiple episodes of carpopedal spasms over a period of next 40 days which were managed by I.V. injection calcium gluconate, and oral calcium and vitamin D₃. She was gradually weaned off the injection, and only oral calcium and vitamin D₃ supplementation were continued. Histopathology was suggestive of Hashimoto’s thyroiditis (Figure 2).

The antithyroid antibody levels were estimated following surgery. Antimicrosomal and antithyroglobulin levels were high. The patient was started on thyroxine 50 mg supplementation and oral vitamin D supplementation was continued.

Thyroid function tests and serum calcium levels were within normal limits at six months following surgery, hence oral calcium was stopped. Thyroxine 50 mg had been continued.

There has been no episodes of hypocalcemic attacks during ensuing three months following stoppage of oral calcium supplementation and serum calcium levels have remained within normal limits.

### DISCUSSION

Grave’s disease and toxic nodular goitre are always considered as the prime etiologies in a patient with thyrotoxicosis. However, certain destructive inflammatory conditions may damage the thyroid gland and cause the classical “leakage” of hormones into the blood resulting in transient thyrotoxicosis [1]. Such phenomenon can occur in Hashimoto’s thyroiditis and called hashitoxicosis.

Hashitoxicosis has an incidence of 4.47%. Out of 69 patients with autoimmune thyroiditis studied by Nabhan et.al. only eight (11.69%) were diagnosed with hashitoxicosis [2]. Normal course of such condition is remission in due course of time or it develops into hypothyroidism.

Certain drugs like pegylated interferons α2b (PEG-IFNα) and ribavirin can produce hashitoxicosis followed by type 1 diabetes [3]. It has multi-factorial etiology which has multiple genetic and environmental factors. Genetic factors include human leucocyte antigen, major histocompatibility complex and cytotoxic T lymphocyte association (CTLA) [4]. Environmental factors include infections, cytokine therapy, selenium, iodine uptake, smoking [5].

It is associated with other autoimmune diseases which includes type 1 diabetes mellitus, systemic lupus erythematosus, multiple sclerosis, rheumatoid arthritis, celiac disease, vitiligo, chronic urticaria [6]. However, our patient did not suffer from any other autoimmune diseases.

In hashitoxicosis, there is a loss of immune tolerance to the thyroid cells due to the production of autoantibodies which in turn leads to destruction of the gland. Hence, the patient presents in hyperthyroid state which is followed by a definite resolution or may go to hypothyroidism. There is a destruction of both follicular and the ‘C’ cells which are replaced by fibrosis [7].

Patients suffering from hashitoxicosis usually present initially with features of hyperthyroidism which later progresses to hypothyroidism [8].

Our patient presented with a short history of nodular goitre and features of toxicity. FNAC revealed colloid material and hence diagnosis of Hashimoto’s thyroiditis was never entertained. The diagnosis of Hashimoto’s thyroiditis can be missed in smears showing cytological evidence of hyperplasia or abundant colloid [9]. Later being the reason in our case. Therefore, it would be reasonable to advocate ultrasonography guided FNAC or multiple punctures in equivocal cases. Macdonald and Yazdi emphasised importance of accurate cytological interpretations in the diagnosis of Hashimoto’s
thyroiditis to avoid false positive and false negative reports [10].

MacDonald and Yezdi emphasised the need for adequate sampling of the thyroid while performing fine-needle aspiration biopsy (FNAB) [11]. In their study of 184 aspirates diagnosed with Hashimoto’s thyroiditis, 39 had corresponding surgical specimen taken from 31 patients. Amongst these 12 (31%) FNABs from nine patients, the cytological diagnosis was not confirmed histologically. The diagnosis of Hashimoto’s thyroiditis is likely to be missed in smears showing cytological evidence of hyperplasia or abundant colloid [11]. In our case too, diagnosis was missed as smears showed only colloid material.

The investigation of choice in cases of hashitoxicosis is the estimation of auto antibodies, which includes antithyroglobulin and antimicrosomal antibodies [1, 11]. Levels of these antibodies are usually raised. The most sensitive of these is the antimicrosomal antibody. In our case, the estimation was done postoperatively and found to be raised. Hence, it will be reasonable to assume that estimation of thyroid antibodies should form integral part of any investigative protocol for thyroid disorders.

Patients with hashitoxicosis are first managed with β blockers to control toxicity and then started with thyroxine supplementation 50 mg once a day in order to slow the disease process and to reduce the level of auto antibodies [11]. However, patients can still undergo surgical resection when there is a dominant mass or an unresponsive mass despite thyroxine therapy or intermediate findings of malignancy on a cutting needle biopsy [12]. Our patient underwent near total thyroidectomy considering it to be a dominant mass of multi-nodular toxic goitre which was not expected to regress.

The patient has been followed-up every six weeks with estimation of thyroid autoantibodies and TSH level. The TSH levels are to be maintained between 1–10 mu/L. If after six months following surgery and thyroxine therapy, autoantibody levels have remained raised, such patients will have to continue life-long thyroxine.

CONCLUSION

In patients with toxic nodular goiter, possibility of Hashimoto’s thyroiditis should be considered and excluded by obtaining representative samples on ultrasonography guided fine-needle aspiration biopsy. Routine estimation of thyroid antibodies should form part of protocol while investigating such patients. There is definite role of surgery under specific clinical settings. Thyroid supplementation will be required for life-long even in those patients who did not undergo any form of surgery.

**********

Author Contributions

Somnath Gooptu – Substantial contributions to conception and design, Analysis and interpretation of data, Drafting the article, Final approval of the version to be published

Gurjit Singh – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Critical revision of the article, Final approval of the version to be published

Iqbal Ali – Acquisition of data, Analysis and interpretation of data, Drafting the article. Final approval of the version to be published

Siddharth Mishra – Acquisition of data, Analysis and interpretation of data, Drafting the article, Final approval of the version to be published

Guarantor

The corresponding author is the guarantor of submission.

Conflict of Interest

Authors declare no conflict of interest.

Copyright

© 2014 Somnath Gooptu et al. This article is distributed under the terms of Creative Commons Attribution License which permits unrestricted use, distribution and reproduction in any medium provided the original author(s) and original publisher are properly credited. Please see the copyright policy on the journal website for more information.

REFERENCES

1. Unnikrishnan AG. Hashitoxicosis: A clinical perspective. Thyroid Research & Practice 2013;10(1):5-6.
2. Nabhan ZM, Kreher NC, Eugster EA. Hashitoxicosis in children: Clinical features & natural history. J Pediatr 2005 Apr;146(4):533-6.
3. Yagyu H, Okada K, Sato S, et al. Pegylated interferon-a2b and ribavirin combination therapy induces Hashitoxicosis followed by type 1 diabetes mellitus. Diabetes Res Clin Pract 2012 Mar;95(3):e52-4.
4. Fisher GF. Molecular Genetics of HLA. Vox Sang 2000;78 Suppl 2:261-4.
5. Boukis MA, Koutras DA, Souvatzoglou A, Evangelopoulou A, Vrontakis M, Moulopoulos SD. Thyroid hormone and immunological studies in endemic goiter. J Clin Endocrinol Metab 1983 Oct;57(4):859-62.
6. Kakourou T, Kanaka-Gantenbein C, Papadopoulou A, Kaloumenou E, Chrousos GP. Increased prevalence of chronic autoimmune (Hashimoto’s) thyroiditis in children and adolescents with vitiligo. J Am Acad Dermatol 2005 Aug;53(2):220-3.
7. Lima MA, Santos BM, Borges MF. Quantitative Analysis of C cells in Hashimoto’s Thyroiditis. Thyroid 1998 Jun;8(6):505-9.
8. Wasniewska M, Corrias A, Salerno M, et al. Outcomes of children with Hashitoxicosis. Horm Res Paediatr 2012;77(1):36-40.
9. Kumar N, Ray C, Jain S. Aspiration cytology of Hashimoto’s thyroiditis in an endemic area. Cytopathology 2002 Feb;13(1):31-9.
10. MacDonald L, Yazdi HM. Fine needle aspiration biopsy of Hashimoto’s thyroiditis. Sources of diagnostic error. Acta Cytol 1999 May-Jun;43(3):400-6.
11. Champion BR, Page KR, Parish N, et al. Identification of a thyroxine-containing self-epitope of thyroglobulin which triggers thyroid autoreactive T cells. J Exp Med 1991 Aug 1;174(2):363-70.
12. Thomas CG Jr, Rutledge RG. Surgical intervention in chronic (Hashimoto’s) thyroiditis. Ann Surg 1981 Jun;193(6):769-6.
Edorium Journals: An introduction

Edorium Journals Team

About Edorium Journals
Edorium Journals is a publisher of high-quality, open access, international scholarly journals covering subjects in basic sciences and clinical specialties and subspecialties.

Invitation for article submission
We sincerely invite you to submit your valuable research for publication to Edorium Journals.

But why should you publish with Edorium Journals?
In less than 10 words - we give you what no one does.

Vision of being the best
We have the vision of making our journals the best and the most authoritative journals in their respective specialties. We are working towards this goal every day of every week of every month of every year.

Exceptional services
We care for you, your work and your time. Our efficient, personalized and courteous services are a testimony to this.

Editorial Review
All manuscripts submitted to Edorium Journals undergo pre-processing review, first editorial review, peer review, second editorial review and finally third editorial review.

Peer Review
All manuscripts submitted to Edorium Journals undergo anonymous, double-blind, external peer review.

Early View version
Early View version of your manuscript will be published in the journal within 72 hours of final acceptance.

Manuscript status
From submission to publication of your article you will get regular updates (minimum six times) about status of your manuscripts directly in your email.

Our Commitment

Six weeks
You will get first decision on your manuscript within six weeks (42 days) of submission. If we fail to honor this by even one day, we will publish your manuscript free of charge.

Four weeks
After we receive page proofs, your manuscript will be published in the journal within four weeks (31 days). If we fail to honor this by even one day, we will publish your manuscript free of charge and refund you the full article publication charges you paid for your manuscript.

Mentored Review Articles (MRA)
Our academic program “Mentored Review Article” (MRA) gives you a unique opportunity to publish papers under mentorship of international faculty. These articles are published free of charges.

Favored Author program
One email is all it takes to become our favored author. You will not only get fee waivers but also get information and insights about scholarly publishing.

Institutional Membership program
Join our Institutional Memberships program and help scholars from your institute make their research accessible to all and save thousands of dollars in fees make their research accessible to all.

Our presence
We have some of the best designed publication formats. Our websites are very user friendly and enable you to do your work very easily with no hassle.

Something more...
We request you to have a look at our website to know more about us and our services.

We welcome you to interact with us, share with us, join us and of course publish with us.

This page is not a part of the published article. This page is an introduction to Edorium Journals and the publication services.