Case Report

Refractory Graves’ Disease Successfully Cured by Adjunctive Cholestyramine and Subsequent Total Thyroidectomy

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The three major forms of treatment for Graves thyrotoxicosis are antithyroid drugs, radioactive iodine therapy and thyroidectomy. Surgery is the definitive treatment for Graves thyrotoxicosis that is generally recommended when other treatments have failed or are contraindicated. Generally, thyrotoxic patients should be euthyroid before surgery to minimize potential complications which usually requires preoperative management with thionamides or inorganic iodine. But several cases of refractory Graves’ disease have shown resistance to conventional treatment. Here we report a 40-year-old female patient with Graves’ disease who complained of thyrotoxic symptoms for 7 months. Her thyroid function test and thyroid autoantibody profiles were consistent with Graves’ disease. One kind of thionamides and β-blocker were started to control her disease. However, she was resistant to nearly all conventional medical therapies, including β-blockers, inorganic iodine, and two thionamides. She experienced hepatotoxicity from the thionamides. What was worse is her past history of serious allergic reaction to corticosteroids, which are often used to help control symptoms. A 2-week regimen of high-dose cholestyramine improved her uncontrolled thyrotoxicosis and subsequent thyroidectomy was successfully performed. In conclusion, cholestyramine could be administered as an effective and safe adjunctive agent for preoperative preparation in patients with severe hyperthyroid Graves’ disease that is resistant to conventional therapies.

Keywords: Graves disease; Drug resistance; Cholestyramine resin

INTRODUCTION

Graves’ disease is the most common cause of thyrotoxicosis. The management of Graves’ disease generally begins with medical therapy [1]. The mainstay of antithyroid medications are the thionamides, such as methimazole (MMI) and propyl-thiouracil (PTU), which usually restore euthyroidism in 1 to 3 months [2].

When the medical treatments have failed, surgery or radioactive iodine (RAI) therapy is generally chosen. If the patient is scheduled to undergo surgery, prompt preoperative achievement of the euthyroid state is recommended to minimize poten-
tial complications. It usually requires pretreatment with thionamides or inorganic iodine. Adjunctive treatment with β-blockers, corticosteroids, and iopanoic acid may also be used to control symptoms [3]. However, literature reviews have reported a few cases of resistant thyrotoxicosis that did not respond to conventional modalities and required additional management [4-6].

Cholestyramine is an ion exchange resin which binds to iodothyronines in the intestines, lowering serum thyroid hormone levels when used adjunctively with thionamides and a β-blocker. Studies from the 1990s have shown cholestyramine to be safe and effective, but its use in clinical practice remains limited [7-10].

In this paper, we report a case of one patient with thyrotoxicosis who was resistant to nearly all conventional medical therapies but successfully responded to high-dose cholestyramine prior to thyroidectomy.

**CASE REPORT**

A 40-year-old female patient presented with symptoms of thyrotoxicosis. She was initially diagnosed with hyperthyroidism 6 years ago and treated with antithyroid medication in local clinics for 5 months. Since then, she maintained a euthyroid state without any medication.

Seven months ago, she started to present thyrotoxic symptoms of fatigue, palpitation, hand tremor, heat intolerance, and dyspnea. Her blood pressure was 110/60 mm Hg, heart rate was 114 beats per minute, and body temperature was 37.2°C. The thyroid gland was diffusely enlarged and firm. Her thyroid function test revealed thyrotoxicosis with elevated thyroid stimulating hormone (TSH) receptor antibody (TRAb); free thyroxine (FT4) of 3.66 ng/dL, triiodothyronine (T3) of 3.65 ng/mL, TSH of 0.08 mIU/L, and TRAb of 4.15 IU/L (normal range: FT4, 0.85 to 1.86; T3, 0.78 to 1.82; TSH, 0.17 to 4.05; TRAb < 1.0). The erythrocyte sedimentation rate (ESR) and C-reactive protein (CRP) level were 47 mm/hr and 0.36 mg/dL, respectively (normal range, ESR, 0 to 15; CRP, 0.01 to 0.47). The electrocardiogram showed sinus tachycardia. The thyroid ultrasonography revealed mild enlargement of both thyroid lobes with heterogeneous echogenicity and no discrete nodules (Fig. 1). The thyroid technetium-99m scan showed right thyroid gland enlargement with inhomogeneously increased activity.

Based on clinical presentation and laboratory findings, the diagnosis of thyrotoxicosis due to Graves’ disease was made. We started MMI and propranolol, both at 30 mg/day, but thyrotoxic signs and symptoms persisted. Despite 4 months treatment, the FT4 and T3 levels increased to 4.27 ng/dL and 7.34 ng/mL, respectively. We increased the dose of MMI up to 45 mg/day but liver enzymes began to increase. Potassium iodide solution 15 mL/day was added for 2 weeks but that did not decrease thyroid hormone levels.

One month after the MMI dose increase, the aspartate aminotransferase and alanine aminotransferase levels had increased to 58 and 101 IU/L, respectively. Therefore, we changed MMI to PTU 300 mg/day and increased the dose of propranolol up to

**Fig. 1.** Thyroid ultrasonography showed mild enlargement of both thyroid lobes with heterogenous echogenicity (A) and increased vascularity (B).
120 mg/day. We asked if she was taking her medication during every visit and she said that she was taking it as recommended. Because she was also taking topiramate, tianeptine, etizolam, and itopride for migraines, we considered the possibility of drug interactions or side effects. Rarely has it been reported to the U.S. Food and Drug Administration that individuals taking these medications developed hyperthyroidism, and there was nothing in the literature that demonstrated a definitive relationship between them.

Despite our efforts, there were no effects on hyperthyroidism and her liver enzymes remained elevated (Table 1). During the treatment, she gained about 12 kg body weight and developed generalized edema.

Refractoriness to antithyroid drugs and inorganic iodides along with the subsequent hepatotoxicity led to the decision to schedule a thyroidectomy. She was admitted to our division of

Table 1. Laboratory Results Change with Drug Challenges before Admission

| Variable | Base | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 |
|----------|------|---|---|---|---|---|---|---|---|
| T3, ng/mL  | 1.33 | 3.65 | 2 | 2.09 | 5.03 | 7.34 | 6.07 | 6.21 | 7.54 |
| FT4, ng/dL  | 1.3 | 3.66 | 2.01 | 2.35 | 4.27 | 3.9 | 3.9 | 3.9 | 3.9 |
| TSH, mIU/L  | 1.26 | 0.08 | 0.07 | 0.04 | 0.11 | 0.06 | 0.08 | 0.12 | 0.08 |
| TRAb, IU/L  | NA | NA | 4.15 | 2.39 | NA | 9.2 | NA | NA | 15.02 |
| AST, IU/L  | 21 | 32 | 20 | 27 | 28 | NA | 60 | 58 | 87 |
| ALT, IU/L  | 16 | 37 | 22 | 37 | 31 | NA | 81 | 101 | 113 |

We started methimazole (MMI) 30 mg/day and propranolol 30 mg/day as initial dose (month 1). Because laboratory data and symptoms suggested the disease was getting worse, doses of MMI and propranolol were gradually increased up to 45 and 120 mg/day, respectively, till 5th month. Subsequently, AST/ALT levels were increased from 28/31 to 60/81 IU/L. After one month treatment, AST/ALT levels were more increased up to 58/101 IU/L so MMI was changed to propylthiouracil 300 mg/day. However, there was no improvement in thyrotoxicosis.

T3, triiodothyronine; FT4, free thyroxine; TSH, thyroid stimulating hormone; TRAb, TSH receptor antibody; NA, not available; AST, aspartate aminotransferase; ALT, alanine aminotransferase.

Fig. 2. Free thyroxine (FT4), total triiodothyronine (T3), and thyroid stimulating hormone (TSH) levels during treatment. The dark gray area in the figure indicates the duration of cholestyramine use until total thyroidectomy. CA, cholestyramine; PTU, propylthiouracil; MMI, methimazole; KI, potassium iodide; PPL, propranolol; HD, hospital day.
endocrinology and metabolism for pre-operative evaluation and close medical management of the thyrotoxicosis. We initially considered glucocorticoids but were not able to use them because she had a history of angioedema, sudden development of skin rash with pruritus, generalized edema, and dyspnea after the intravenous administration of steroids. A few cases in which high-dose cholestyramine successfully treated refractory thyrotoxicosis had been reported [5], so we added 12 g divided three times a day to the patient’s treatment regimen.

The patient’s T3 levels began to decrease dramatically and there was a notable improvement of symptoms (Fig. 2). She had a robotic total thyroidectomy without any complications (Fig. 3).

**DISCUSSION**

There are several cases in the literature of refractory Graves’ disease that resistant to conventional therapies. Refractory cases have shown resistance to high-dose thionamides, β-blockers, and rarely, to iodine [4-6,11-13]. In an attempt to restore euthyroidism in refractory cases before surgery, several modalities have been reported. Iopanoic acid, an oral cholecystographic agent, was used for rapid preoperative preparation of refractory thyrotoxicosis had been reported [5], so we added 12 g divided three times a day to the patient’s treatment regimen.

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In our patient, resistance to two different thionamides (MMI and PTU) and iodine was observed. Also, propranolol did not improve the symptoms of thyrotoxicosis. Malabsorption was ruled out by clinical and biochemical data. Serum and thyroid MMI levels and the presence of antidrug antibodies were not obtained in this case, but rapid drug metabolism, the existence of antidrug antibodies and impaired accumulation of MMI must be considered as possible reasons for drug resistance.

However, the most probable reason for refractoriness to antithyroid medications in this case is the severity of the hyperthyroidism, particularly high T3 level. Dahlberg et al. [14] found that the failure of carbimazole in the treatment of hyperthyroidism is primarily dependent upon the T3 level. When the T3 level is above 6 ng/mL before treatment, a higher dose of antithyroid drug was needed. Because the increase of liver enzyme was mild in our patient, we could not completely rule out the possibility of uncontrolled thyrotoxicosis as the reason of the liver enzyme abnormality. But we could not administer an unconventionally high-dose of MMI because of the risk of hepatotoxicity. During preparation for surgery, we also had limitation to use of glucocorticoids, because of the possibility of anaphylactic reaction from her past history. Rarely patients with hypersensitivity reactions to corticosteroids were reported [15]. Clinical manifestations of steroid hypersensitivity range from relatively minor reactions such as urticaria to more life-threatening ones, such as bronchospasm and anaphylaxis. Studies have shown that the most common steroids to which people develop hypersensitivity reactions to are hydrocortisone and methylprednisolone. A low dose challenge with oral prednisolone was considered, but the patient refused.

Cholestyramine is an anion exchange resin which binds bile in the gastrointestinal tract to prevent its reabsorption. It is mainly used to lower serum cholesterol and to treat pruritus.
The known primary side effects of cholestyramine are constipation and abdominal discomfort. However, the aforementioned studies showed only a few minor complications like hyperdefecation and bad taste. Our patient also tolerated cholestyramine without any complications. Remarkably, in Tsai’s study [9], which only included Chinese individuals, no one experienced constipation. They suggested high fiber diets in Chinese culture as the possible reason. Also considering that the main gastrointestinal symptom of thyrotoxicosis is diarrhea, constipation as a side effect may be favorable.

In conclusion, we recommend cholestyramine as an effective adjunctive therapy for preoperative preparation in patients with severe Graves’ disease that is resistant to conventional therapies or when a failure of conventional therapies is expected due to high T3 level. Considering the safety profiles and dramatic effects of cholestyramine, it may be the primary option for treatment of severe thyrotoxic patients, along with antithyroid drugs. Also in contrast to antithyroid drugs which inhibit thyroid hormone synthesis and are used for the hyperthyroidism, cholestyramine could be an option for treatment of any condition with excessive thyroid hormone levels, including thyroiditis and factitious thyroid hormone disorder [16].

CONFLICTS OF INTEREST
No potential conflict of interest relevant to this article was reported.

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