Thyrotoxic Pericardial Effusion Complicating Graves’ Disease in Pregnancy

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Abstract

Pericardial effusion is a rare complication of Graves’ disease. A pregnant Filipino woman with diffuse goiter and hyperthyroid symptoms was initially treated as a case of Graves’ disease. She was readmitted for exertional dyspnea, orthopnea, and bipedal edema; an echocardiogram revealed a massive pericardial effusion. Workup for other causes of the effusion was unremarkable. The pericardial effusion resolved after pericardiostomy and anti-thyroid medications. This highlights the clinician’s role in determining the association between the two disease entities.

Keywords: Graves’ disease, thyrotoxicosis, hyperthyroidism, pericardial effusion, pregnancy

BACKGROUND

While sinus tachycardia, atrial fibrillation, and congestive heart failure are well-recognized cardiovascular manifestations of Graves’ disease,1 pericardial effusion due to thyrotoxicosis is rarely reported. This case highlights the clinician’s role in determining the association between the two disease entities.

CASE

A 32-year-old Filipino woman on the 19th week of her fourth pregnancy presented at the emergency room with a seven-year history of diffuse goiter associated with tremors, palpitations, heat intolerance, exophthalmos and exertional dyspnea. Prior to this consult, she was previously taking anti-thyroid medications which were discontinued five years ago upon improvement of symptoms, and was subsequently lost to follow up. On admission, the patient was febrile, tachycardic in atrial fibrillation, hyperreflexic with fine finger tremors. Ophthalmologic examination revealed positive Dalrymple’s, Kocher’s, and Von Graefe’s signs, lagophthalmos, chemosis and periorbital edema. The thyroid gland was non-tender and diffusely enlarged, measuring 5x5 cm per lobe. Auscultation revealed bibasal fine crackles, and there was note of hyperpigmented, non-pitting in duration over the pretilial area of both legs, clubbing of the fingers and pedal edema. She was treated for Graves’ disease in thyroid storm, with congestive heart failure from thyrotoxic heart disease. Her chest radiograph showed left ventricular cardiomegaly (Figure 1). She improved with propylthiouracil, propranolol and furosemide and was discharged.

Figure 1. Chest radiograph taken on postero-anterior view during initial admission showing left ventricular cardiomegaly with pulmonary congestive changes.
Nineteen days later, the patient noted recurrence of exertional dyspnea, bipedal edema and three-pillow orthopnea. No chest pain or fever was observed. Upon readmission, she was tachycardic with distented neck veins, muffled heart sounds and Grade 2 bipedal edema. Deep tendon reflexes were normal with no tremors. Chest radiograph now revealed multichambered cardiomegaly (Figure 2.)

![Chest radiograph on readmission showing multichambered cardiomegaly and characteristic water-bottle sign indicative of massive pericardial effusion.](image)

**Figure 2.** Chest radiograph on readmission showing multichambered cardiomegaly and characteristic water-bottle sign indicative of massive pericardial effusion.

**Laboratory Workup**

During the patient’s first admission, free thyroxine (FT4) levels were elevated (40.2 pmol/L), and thyroid stimulating hormone (TSH) was low (0.1 uIU/mL). On readmission, FT4 levels were normal (11.8 pmol/L) while TSH remained low (0.005 uIU/mL).

Her 12-lead electrocardiogram showed atrial fibrillation in rapid ventricular response, with nonspecific ST-T wave changes. A 2D echocardiogram with color-flow Doppler during the second admission showed normal-sized chambers with left ventricular remodelling and mildly depressed left ventricular systolic function; adequate right ventricular systolic function; mild to moderate pulmonary hypertension; and massive pericardial effusion in impending tamponade (Figure 3). Pericardial fluid samples were sent for differential cell counts, cytology, acid-fast (AFB) and conventional gram stain and culture, tuberculous polymerase chain reaction (TB PCR) and determination of glucose and total protein levels. Table 1 summarizes the pericardial fluid analysis.

**Table 1. Pericardial Fluid Studies**

<table>
<thead>
<tr>
<th>Qualitative Examinations</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Color</td>
<td>Dark red</td>
</tr>
<tr>
<td>Transparency</td>
<td>Slightly hazy</td>
</tr>
<tr>
<td>Red Blood Cells</td>
<td>283,000 x 10⁹/L</td>
</tr>
<tr>
<td>White Blood Cells</td>
<td>578 x 10⁹/L</td>
</tr>
<tr>
<td>Polymorphonuclear Cells</td>
<td>82%</td>
</tr>
<tr>
<td>Lymphocytes</td>
<td>17%</td>
</tr>
<tr>
<td>Distorted Cells</td>
<td>1%</td>
</tr>
<tr>
<td>Glucose</td>
<td>4.95 mmol/L</td>
</tr>
<tr>
<td>Total Protein</td>
<td>59.01 mg/dL</td>
</tr>
<tr>
<td>Gram Stain</td>
<td>PMN 0-1/OIF</td>
</tr>
<tr>
<td>Bacterial Culture</td>
<td>No growth</td>
</tr>
<tr>
<td>AFB Smear</td>
<td>Negative</td>
</tr>
<tr>
<td>AFB Culture</td>
<td>No growth</td>
</tr>
<tr>
<td>TB PCR</td>
<td>Negative</td>
</tr>
<tr>
<td>Cell Block and Cytology</td>
<td>Negative</td>
</tr>
</tbody>
</table>

Fetal screening with congenital anomaly scan showed no gross fetal structural abnormalities, with an estimated sonographic weight that was appropriate for gestational age. Serum antinuclear antibody (ANA) to rule out the possibility of an autoimmune process was likewise negative.

**Treatment**

The patient underwent pericardiocentesis with low flow drainage while on dobutamine infusion, draining around 600 cc of serosanguinous fluid. Dobutamine was discontinued after the procedure, and the patient’s vital signs remained stable throughout admission except for intermittent episodes of tachycardia. Medical therapy included methimazole, digoxin, prednisone and furosemide. Propranolol was also initiated after initial diuresis to control the heart rate and decrease the adrenergic effects of the hyperthyroid state. The patient...
experienced gradual resolution of orthopnea, pedal edema and exertional dyspnea, with spontaneous conversion to normal sinus rhythm. On the 18th hospital day, she was subsequently discharged.

Outcome and Follow up

Two months after discharge, the patient underwent a repeat 2D echocardiogram (Figure 4) revealing the absence of the pericardial effusion but with pericardial thickening. There was also improvement of overall left ventricular systolic function after treatment with heart failure and anti-thyroid drugs. The patient went into preterm labor at 32 weeks of gestation. She delivered a live baby girl via spontaneous vaginal delivery, who was admitted to the neonatal intensive care unit and eventually discharged.

Figure 4. Repeat 2D echocardiogram after two months, showing absence of pericardial effusion but with thickened pericardium.

DISCUSSION

Our patient initially presented with classic thyrotoxic symptoms associated with diffuse goiter, typical eye signs, pretibial dermopathy and acropathy. These findings, plus a high FT4 and low TSH, make the clinical diagnosis of Graves’ disease straightforward. In thyrotoxic patients, the most common cardiac complications are sinus tachycardia, atrial fibrillation and congestive heart failure, with reports of heart block and a cardiomyopathy that may or may not be reversible following return to a euthyroid state. These manifestations may be exaggerated in pregnancy owing to the physiologic increase in heart rate, stroke volume and cardiac output.

In thyroid disorders, pericardial effusion is more commonly seen in 5-30% of hypothyroid patients. In contrast, its occurrence in Graves’ disease is somewhat rare, one study looking into the causes of pericardial effusion in 204 patients found no cases of thyrotoxicosis among them. This lack of published evidence linking pericardial effusion with thyrotoxicosis emphasizes the importance of a high index of suspicion to ascertain the association between the two disease entities.

Our initial consideration for the patient’s effusion was an infectious etiology, specifically Mycobacterium tuberculosis, which is highly prevalent in the Philippines. Given that the patient is a female of reproductive age, the possibility of autoimmune pericarditis secondary to an underlying connective tissue disease such as systemic lupus erythematosus was also entertained. Malignancy was considered since it was found to be the most common etiology in a study among Filipino patients with pericardial effusion in tamponade. We also regarded the possibility that the effusion may have been due to the pregnancy itself, but it is usually mild and characteristically manifests as a transudate. Finally, the effusion may have been part of an unusual vasculitic side-effect from PTU; however, this was also unlikely as the other typical vasculitic manifestations of rash, hemoptysis, cytopenia, hematura and renal failure were absent in the patient. Since workup for all these causes eventually turned out unremarkable (the gram stain result was most likely a contaminant), with the effusion resolving after pericardiostomy and anti-thyroid drugs (making an idiopathic cause less likely), we concluded that thyrotoxicosis caused the patient’s effusion.

It is interesting that the patient’s effusion was detected during readmission when she had already received antithyroid drugs and FT4 levels had normalized. This is consistent with reports showing that pericardial effusion resulting in tamponade can still develop in Graves’ thyrotoxicosis even during anti-thyroid treatment and with improving thyroid function tests. Moreover, although the patient was initially discharged improved, the fact that some of her hyperthyroid symptoms (tachycardia, dyspnea, edema) persisted on readmission with TSH being suppressed further from 0.1 to 0.005 uIU/mL showed that the thyrotoxicosis was still not adequately controlled. PTU, initially started due to its generally better safety profile for pregnancy, was subsequently shifted to methimazole, a more potent antithyroid drug.

Since the apparent worsening of the patient’s heart failure symptoms on readmission was temporally associated with her recent thyrotoxic manifestations, thyrotoxicosis was considered as the underlying cause of the heart failure, despite the initiation of standard cardiac medications. This is consistent with reports showing that in some patients treated for hyperthyroidism, achievement of a euthyroid state is not by itself sufficient to fully reverse cardiomyopathy and left ventricular failure. Furthermore, the patient’s pregnant condition, persistent tachycardia, and atrial fibrillation could have also contributed independently to the exaggeration of the heart failure symptoms.
The pathophysiology of thyrotoxic pericardial effusion has been demonstrated to be similar to that of pretilial myxedema, with a characteristic serous effusion attributed to the transudation of albumin and decreased lymphatic clearance of interstitial fluid proteins. On occasion, the effusion can also appear serosanguinous, as in our patient. This purported underlying pathophysiologic mechanism is the basis for several reports stating the potential benefit of prednisone in these patients. Hence, we decided to continue prednisone even after TB was ruled out as an unlikely etiology of the effusion.

Even with treatment, the mortality rate for thyrotoxicosis from cardiac sequelae can reach 30%. An echocardiogram is therefore recommended as part of the investigation to rule out pericardial effusion as a contributing factor potentially heralding the development of cardiac tamponade. Conversely, in a patient presenting with an unexplained pericardial effusion, it might be prudent to exclude Graves’ thyrotoxicosis despite the absence of classic signs and symptoms, in addition to routine tests for microbiology, inflammatory and oncologic markers and autoimmunity.

This is the first documented case of its kind in the Philippines, and in the entire ASEAN region. Moreover, in contrast to other reports on thyrotoxicosis-related pericardial effusion, this is also so far the first case dealing with a pregnant patient. The rarity of the association emphasizes the need for vigilance.

References