DESCRIPTION of CASE

A 35-year-old woman became aware of a swelling in her neck about ten weeks before referral. She had been trying to conceive for ten months. She had had a missed abortion five months earlier. Because of the neck swelling, her family doctor arranged thyroid function tests, which were in the thyrotoxic range on two occasions five weeks apart: serum free thyroxine, 26 and 28 pmol/l (normal range, 11–23); serum free tri-iodothyronine, 10.9 and 11 pmol/l (normal range, 3.5–6.5); serum thyroid-stimulating hormone (TSH), <0.05 mU/l (normal range, 0.3–4.1).

Her previous medical history included a partial thyroidectomy for thyrotoxicosis at the age of 24. Other than a goitre, she had no symptoms except increased appetite and a slight tremor, which she had been aware of for about eight weeks. Following the missed abortion, she had two normal menstrual periods. Her only medication was folic acid supplements. She worked part-time and had a two-and-a-half-year-old child.

On examination she was of average weight. Her hands were warm and moist. There was a fine tremor. A previous thyroidectomy scar was noted. The right lobe of the thyroid was palpable and felt smooth. There was a bruit over the right thyroid lobe on auscultation. She had lid retraction and lid lag but no other signs suggestive of thyroid-associated ophthalmopathy (TAO) (Figure 1). Her pulse rate was 100 beats per minute and regular. Her blood pressure was 150/70 mm Hg. The rest of the examination was normal.

What Is the Cause of Her Thyrotoxicosis?

Thyrotoxicosis is no more than a descriptor for a pattern of biochemical abnormalities. Before considering treatment, it is the clinician’s task to define the underlying cause, as an accurate diagnosis is an essential guide to the most appropriate treatment (Box 1).

The most likely causes in this case were Graves disease, thyroiditis, toxic multinodular goitre (TMNG), and toxic adenoma. The hallmark of TMNG or toxic adenoma is the presence of one or more palpable thyroid nodules. In this case the patient had previously undergone a partial thyroidectomy and a vascular thyroid remnant was palpable on the right thyroid lobe. Post-partum thyroiditis occurs within 12 months of childbirth; a variant of this condition occurs after miscarriage. In this patient’s case post-partum thyroiditis was unlikely because her previous pregnancy was 2.5 years earlier; however, the miscarriage five months earlier may have been relevant. Viral thyroiditis is usually preceded by an upper respiratory tract infection and the thyroid gland is tender to touch; the absence of these features makes viral thyroiditis unlikely. “Silent” thyroiditis may present in this way and was a possibility here.

Laboratory tests that may help differentiate between the different causes of thyrotoxicosis include a radiolabelled technetium or iodide thyroid scan (Figure 2), and measurement of anti–thyroid peroxidase (TPO) antibodies, TSH receptor antibodies, and inflammatory markers (Table 1). The thyrotoxic phase of thyroiditis is usually followed by spontaneous euthyroidism and in some cases hypothyroidism. Repeating thyroid function tests within a few weeks of the first set may identify cases of thyroiditis.

In this case the prolonged time course of thyrotoxicosis, the presence of a vascular thyroid remnant, the persistently thyrotoxic thyroid function tests, and the elevated serum

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Abbreviations: 131I, radiiodine; TAO, thyroid-associated ophthalmopathy; TMNG, toxic multinodular goitre; TPO, thyroid peroxidase; TSH, thyroid-stimulating hormone

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levels of TSH receptor antibodies (62 U/l; reference range, 0–10) were in favour of a diagnosis of recurrent Graves disease.

What Are the Effects of Thyrotoxicosis on Fertility and Risk of Abortion?

Menstrual irregularities occur in about 20% of thyrotoxic women [1]. Infertility is common in women with thyrotoxicosis even when they maintain ovulatory cycles [1]. Thyrotoxicosis also increases the risk of miscarriage to 26% [2].

How Should This Patient Be Treated?

There are three treatment options for thyrotoxicosis due to Graves disease: radioiodine (131I) therapy, thyroidectomy, and anti-thyroid drugs [3]. 131I therapy is safe and effective, but pregnancy should be deferred for 4–6 months after treatment as there are theoretical risks of fetal abnormalities. Most national regulatory authorities recommend avoidance of close contact with adults for a few days and with children and pregnant women for 2–3 weeks. 131I therapy was not appropriate for this patient because she wished to proceed with pregnancy as soon as possible and she had a two-and-a-half-year-old child, who would be difficult to care for after 131I therapy.

A second thyroidectomy is worthy of consideration, but involves general anaesthesia and a period of recuperation of a few weeks and therefore disruption of family and professional life. The risks of damage to the recurrent laryngeal nerves and parathyroid glands after a second thyroidectomy are considerably greater than after a first operation and are of the order of 5%–10%. Because of these considerations, thyroidectomy was not felt to be a suitable option.

Anti-thyroid drugs (carbimazole, methimazole, and propylthiouracil) restore euthyroidism within a few weeks of initiation of treatment [4]. Minor side effects (such as skin rashes) occur in about 5% of cases. Agranulocytosis is rare (~0.4%), but the consequences are life threatening and all patients on anti-thyroid drugs must be made aware of this complication (Box 2). All anti-thyroid drugs have been used and are acceptable in pregnancy.

Congenital anomalies have been reported in association with anti-thyroid drugs, but the increase in risk above background is very marginal. The risks of aplasia cutis and choanal and oesophageal atresia may be slightly lower with propylthiouracil than with other anti-thyroid drugs (choanal and oesophageal atresia, scalp defects, minor facial anomalies, and psychomotor delay compose an embryopathy implicated with methimazole use). But because the evidence is inconclusive and the additional risk minimal, all three drugs are widely used in pregnancy. The lowest dose of anti-thyroid drug that maintains euthyroidism should be used in women who wish to become or are already pregnant, in order to avoid fetal hypothyroidism and fetal goitre formation.

In this case propylthiouracil was used initially, at a dose of 50 mg four times per day. The patient was advised to take contraceptive measures until euthyroidism. Four weeks later her thyroid function tests had improved: serum free thyroxine, 13 pmol/l; serum total tri-iodothyronine, 2.5 nmol/l (normal range, 1.34–2.73); serum TSH, <0.05 mU/l. The dose of propylthiouracil was reduced to 25 mg four times per day, and the patient was advised that she could start trying to conceive.

What Are the Risks of TAO?

TAO is a complication that many patients fear. It can be disfiguring and difficult to treat [3]. If there are no clinical features of TAO at presentation, the risk of developing it in future is approximately 15%. Smoking is an important predisposing factor. As this patient was a non-smoker the probability of developing TAO is less than 10%.

What Monitoring Is Required during Pregnancy?

The dose of anti-thyroid drug usually needs to be decreased during pregnancy, and often Graves disease remits completely.

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**Table 1. Diagnostic Tests for Identifying the Cause of Thyrotoxicosis**

| Condition                  | Technetium 99 or Iodide Scan | Anti-TPO Antibodies | Anti-TSH Receptor Antibodies | Inflammatory Markers (ESR/CRP) |
|----------------------------|------------------------------|---------------------|-----------------------------|-------------------------------|
| Graves disease             | Increased uniform uptake     | 70% positive        | >95% positive               | Normal                        |
| TMNG                       | “Hot” and “cold” areas       | 0%–5% positive      | <5% positive                | Normal                        |
| Toxic adenoma              | Single “hot” area            | 0%–5% positive      | <5% positive                | Normal                        |
| Thyroiditis—viral          | Decreased uptake             | 0%–5% positive      | <5% positive                | Elevated                      |
| Thyroiditis—silent/post-partum | Decreased uptake         | >90% positive       | <5% positive                | Normal                        |

CRP, C-reactive protein; ESR, erythrocyte sedimentation rate.
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and the medication can be withdrawn. This is probably due to
the overall immunosuppressive effect of pregnancy.

Monitoring of free thyroid hormone concentrations is of
paramount importance during pregnancy and should be
performed every 4–6 weeks, or more frequently if thyroid
status is changing. The biochemical target is to achieve and
maintain maternal serum free thyroxine levels at or slightly
above the upper limit of normal, using the lowest dose of
anti-thyroid drug possible. TSH receptor antibodies should be
measured in the third trimester because positivity is predictive
of neonatal thyrotoxicosis [5].

When the mother (as in this case) has a functioning
thyroid gland or remnant in situ, maternal thyroid function
mirrors that of the fetus. If there are concerns about fetal
thyrotoxicosis (e.g., because maternal hyperthyroidism
proves difficult to control), fetal heart rate monitoring
should be undertaken. A persistent fetal tachycardia
greater than 160 beats per minute is suggestive of fetal
thyrotoxicosis. In cases where fetal thyrotoxicosis is
diagnosed, monitoring of fetal growth and fetal goitre by
ultrasound is imperative. In most cases the fetus can be
treated satisfactorily by adjusting the dose of anti-thyroid
drug in the mother and by following the fetal response
clinically and by ultrasound.

What Are the Risks to the Fetus in a Woman with
Graves Disease?

Poor control of maternal hyperthyroidism is associated with
significant obstetric complications including miscarriage
(26%), low birth weight, prematurity, (pre-)eclampsia, and
possibly congenital malformations [6]. After the fetal thyroid

Box 2. Patient Information Leaflet Used
by the Author to Remind Patients Receiving
Anti-Thyroid Drugs of the Potential Complication
of Agranulocytosis

“You have been started on a drug called Carbimazole/
Methimazole/Propylthiouracil to control the activity of your
thyroid gland. This is important treatment and Carbimazole/
Methimazole/Propylthiouracil is a well established drug that
has been used for many years. The great majority of people
treated with Carbimazole/Methimazole/Propylthiouracil have no
problems whatsoever.

“Some people occasionally develop a rash—if this happens
please consult your doctor as soon as possible; you need not
discontinue the drug unless he/she tells you to do so.

“More rarely, Carbimazole/Methimazole/Propylthiouracil
affects white cells in the blood, in which case you would be likely
to develop a very severe sore throat and to feel ill with a fever.
If this happens while you are on Carbimazole/Methimazole/
Propylthiouracil treatment you must attend either your family
doctor or the hospital on the same day, to have your blood count
checked. Take no more tablets until the blood count has been
checked. If your white blood count is normal you can carry on
with the Carbimazole/Methimazole/Propylthiouracil. If your
white blood count is abnormal your family doctor or the hospital
will need to deal with this problem urgently.

“Please keep this with you in case you need to show it to your
doctor.”

maters (from 20 weeks of gestation onwards), maternal TSH
receptor antibodies may act on the fetal thyroid to cause fetal
thyrotoxicosis and goitre. The risk of fetal thyrotoxicosis is
about 1% of all pregnancies in women with Graves disease,
and if untreated, fetal mortality may be as high as 24%.
Overtreatment may lead to hypothyroidism in the fetus,
which is associated with subtle neurocognitive deficits later
on in life, particularly if the hypothyroidism occurs in the
first trimester [7]. Fetal goitre can develop as a result of fetal
thyrotoxicosis or fetal hypothyroidism and in severe cases can
obstruct labour.

What Are the Risks of Recurrence of Thyrotoxicosis
after Delivery?

The risk of relapse of maternal thyrotoxicosis is high in
the post-partum period (up to 80%), and close monitoring
is required. Anti-thyroid drugs can be used safely during
breastfeeding [8].

Prenatal Counselling of Women with Graves Disease

Pregnancy is a common concern among women of
childbearing age who are receiving treatment for Graves
disease. Some women may elect to have definitive treatment
before pregnancy, which can be either a thyroidectomy or
131I therapy. The advantage of these treatment options is
that the risk of maternal thyrotoxicosis during pregnancy
is reduced, if not eliminated. Fertility is not affected by
131I therapy for thyrotoxicosis, but pregnancy should be
deferred for 4–6 months after 131I therapy, although the
basis of this recommendation is largely empirical. The risk
of fetal and neonatal thyrotoxicosis is not eliminated by
previous thyroidectomy or 131I therapy. The most important
treatment for Graves disease is to work with their practitioner to ensure that
thyroid function tests are normal at the time of conception and
throughout pregnancy.
Investigating the Cause of Thyrotoxicosis

In many cases of thyrotoxicosis the aetiology will be apparent from information that can be obtained from the history and clinical examination. In cases where there is doubt, additional investigations are indicated. Direct measurement of TSH receptor antibody levels is not widely available, but can be very valuable as modern assays are highly sensitive and specific. TSH receptor antibodies can occasionally be positive in post-partum thyroiditis (this seems to be particularly rare in Europe, though reported in North America and Japan), and in cases of doubt a thyroid scan showing no uptake of radioisotope is diagnostic of thyroiditis [9]. TSH receptor antibody measurement is indicated in pregnancy to assess the risks of fetal and neonatal thyrotoxicosis. Anti-TPO antibodies occur in a significant proportion of the normal population, and this limits the use of this test. High concentrations of anti-TPO antibodies are present in silent and post-partum thyroiditis. Radioisotope scans are useful in identifying the cause of thyrotoxicosis (Figure 2), but should be avoided in pregnancy.

Learning Points
- Thyrotoxicosis is not a diagnosis, merely a biochemical result. An accurate clinical diagnosis encompassing the aetiology is imperative for optimal management.
- The most common cause of thyrotoxicosis in women of childbearing age is Graves disease.
- Thyrotoxicosis impairs fertility, and thyroid status should be assessed in women with secondary infertility or recurrent abortions.
- Three treatments are available for thyrotoxicosis due to Graves disease: anti-thyroid drugs, \(^{131}I\) therapy, and thyroidectomy. The right treatment is that which suits the patient’s individual circumstances best.
- \(^{131}I\) therapy is an absolute contraindication in pregnancy. Anti-thyroid drugs may be used safely, and the dose should be titrated to the minimum dose that maintains normal maternal thyroid hormone levels.
- The hyperthyroidism of Graves disease usually remits after the first trimester, and anti-thyroid drugs can be withdrawn; however, relapse of maternal thyrotoxicosis in the post-partum period is common.
- Uncontrolled maternal hyperthyroidism can lead to fetal thyrotoxicosis with devastating effects on the fetus. Fetal thyrotoxicosis can be treated satisfactorily by appropriate manipulation of the maternal dose of anti-thyroid drug and careful fetal monitoring.
immunosuppressive effect of anti-thyroid drugs on synthesis of TSH receptor antibodies, but relapse is the rule in cases of TMNG or toxic adenoma.  

$^{131}$I therapy is effective for Graves disease, TMNG, and toxic adenoma. $^{131}$I therapy is ineffective in thyroiditis because iodine uptake is reduced or absent in this condition (Figure 2). Most patients with Graves disease develop permanent hypothyroidism after $^{131}$I therapy, whereas most patients with TMNG and toxic adenoma do not. $^{131}$I therapy is associated with a small risk of exacerbation of new development of TAO, particularly in smokers.  

Thyroiditis may require symptomatic treatment with beta blockers during the thyrotoxic phase.

The type of thyroidectomy (subtotal versus total) for Graves disease as primary treatment has been the subject of controversy for some years. The argument in favour of total thyroidectomy is that the risk of recurrence of the thyrotoxicosis is eliminated, and that if performed by skilled thyroid surgeons the probability of hypoparathyroidism and vocal cord palsy is no greater than for a subtotal thyroidectomy [10]. A subtotal thyroidectomy, on the other hand, provides the best chance of any treatment for Graves disease for long-term euthyroidism without the need for thyroxine or other treatments for thyrotoxicosis.

The choice of treatment for Graves disease should be tailored to the needs of the individual patient, but also depends on local facilities, surgical expertise, and patient choice.

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