Thyroid dysfunction in pregnancy

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Pregnancy induces complex hormonal and immunological changes that modify normal thyroid physiology. Therefore, evaluation of thyroid function during pregnancy should be interpreted according to these changes. In our opinion, the high prevalence of pregnancy-related thyroid disorders and their important consequences for both mother and fetus indicate the need for routine thyroid function screening both before and during pregnancy. Once thyroid dysfunction is diagnosed, the management of the disorder requires frequent monitoring to adjust treatment accurately. The goal of treating hyperthyroidism with thionamide drugs is to maintain serum thyroxin (T4) in the upper normal range (free T4, 2-2.5 ng/dl; total T4, 12.0-18.0 µg/dl) using the lowest possible dose of the drug, while in hypothyroidism the goal is to return serum thyrotropin to the range between 0.5 and 2.5 mU/l.


INTRODUCTION

Pregnancy is a unique situation in which the physician is faced with at least two interactive patients. Any medical action or inaction may have positive or negative consequences for both the mother and the fetus.

Maternal physiological changes during pregnancy

The physiological changes that occur in normal pregnancy have important repercussions for the thyroid gland. Usually thyroid gland volume enlarges and thyroid hormone production increases approximately 50% above the preconception baseline. These changes are secondary to a variety of factors.

Human chorionic gonadotrophin. Rising plasma levels of placental human chorionic gonadotrophin (hCG), which has a weak thyrotropin (TSH) agonist action due to the structural homology

Revisiones

DISFUNCIÓN TIROIDEA EN EL EMBARAZO

El embarazo comporta una serie de cambios hormonales e inmunológicos que dan lugar a modificaciones en la fisiología normal del tiroides. Por tanto, la evaluación de la función tiroidea durante el embarazo debe interpretarse teniendo en cuenta estos cambios. En nuestra opinión, la gran prevalencia de trastornos tiroideos asociados al embarazo y las graves consecuencias que pueden causar a la madre y el feto apoyan la necesidad de realizar pruebas de cribado de disfunción tiroidea de manera sistemática, tanto antes como durante el embarazo. Una vez se ha diagnosticado una disfunción tiroidea será necesario realizar una monitorización frecuente para ajustar el tratamiento de forma precisa. El objetivo del tratamiento del hipertiroidismo con fármacos antitiroideos es lograr que la concentración de tiroxina sérica (T4) se mantenga en el límite alto del rango normal (T4 libre, 2-2.5 ng/dl; T4 total, 12-18 μg/dl) con la mínima dosis posible, mientras que, en el caso del hipotiroidismo, el objetivo es conseguir que la concentración de tirotropina se mantenga entre 0.5 y 2.5 mU/l.

between both hormones, have a major thyroid stimulatory influence. hCG increases during the first trimester and plateaus from midgestation to shortly after delivery. The result of this hCG activity is an elevation in serum thyroxine (T4) and triiodothyronine (T3) concentrations and suppression of serum TSH, providing new normal ranges unique to pregnancy (fig. 1).

Thyroid-binding globulin. Plasma oestrogen levels rise in pregnancy and induce an elevation of up to 100% in serum thyroid-binding globulin (TBG). This occurs mainly during the first 20 weeks and is secondary to an extended half-life because of changes in TBG glycosylation. As a result, by approximately week 10 of pregnancy the total serum T4 (TT4) is elevated by up to 50%, remaining constant at this level until delivery. This large increase in TBG opens many T4 binding sites, which have to be filled to maintain free T4 equilibrium and, therefore, constitute another cause of increased thyroid hormone secretion. These ongoing changes in TBG have made assessment of free thyroid hormone levels in pregnancy a technical challenge, resulting in a polluted literature in which cross-sectional studies have suggested that free T4 (FT4) levels during the first trimester may be higher, lower or the same as those before conception (fig. 2).

Other important factors influencing thyroid function. During pregnancy, other physiological adjustments take place in maternal thyroid homeostasis which, together, may lead to incremental increases in thyroid hormone synthesis. The maternal glomerular filtration rate is also elevated secondary to increased cardiac output, resulting in high renal clearance and iodide excretion. Therefore, iodine intake needs to be increased to accommodate the continuing thyroid hormone synthesis. In addition, transplacental passage of T4 may also stimulate the maternal thyroid by depleting maternal circulating T4.

Immune changes. Pregnancy is a time of placenta-induced immune suppression secondary to placental cytokine and hormone secretion resulting in enhanced regulatory T cell function. This particular situation can be extremely important to autoimmune reactions and most autoimmune diseases, including thyroid disorders, tend to improve during gestation. Maternal thyroid function is totally reset to normal activity by 6 months after delivery unless thyroid dysfunction develops.

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Fig. 1. Thyroid-stimulating hormone (TSH) plasma concentration expressed in percentiles according to gestational age. TSH was measured in 13,599 singleton pregnancies. Gestational age-specific. Modified from Dashe et al. 33

Fig. 2. Changes in plasma concentrations of thyroid function tests and hCG according to the evolution of pregnancy. The shaded area corresponds to the normal range in non-pregnant women. hCG: human chorionic gonadotrophin; TBG: thyroid-binding globulin; T4: thyroxine; TSH: thyroid-stimulating hormone. Modified from Brent GA. Maternal thyroid function: interpretation of thyroid function tests in pregnancy. Clin Obstet Gynecol. 1997;40:3-15.
Fetal thyroid physiology

The fetal thyroid starts producing sufficient hormones for the fetus by the end of the first trimester. Before this time, the fetus is dependent on a supply of maternal thyroid hormones, which is metabolically controlled by the placental deiodinase enzymes. The placenta expresses all three deiodinases D1, D2 and D3. D2 converts the pro-hormone T4 into the biologically active T3, whereas D3 inactivates T3. The role of D1 is less important. D3 is by far the most prevalent isoform in the placenta and consequently fetal T3 plasma concentrations are low until the 30th week of gestation.

Maternal thyroid function, especially in the first trimester, is, therefore, extremely important to adequate fetal central nervous system (CNS) development. Recent studies have shown that there are specific thyroid hormone transporters in the fetal CNS that play an important role in fetal CNS development, but their consideration is beyond the scope of this review.

ASSESSMENT OF THYROID FUNCTION DURING PREGNANCY

Gestation is a prime time for the diagnosis of thyroid diseases, insofar as gravid women usually seek regular medical care. Indeed, thyroid diseases are the most common group of pre-gestational endocrine diseases that persist during pregnancy. Although thyroid dysfunction is an important barrier to pregnancy, significant thyroid dysfunction still occurs in 1-2% of pregnant women and mild forms of thyroid disease are even more prevalent. The presence of either hyperthyroidism or hypothyroidism as evidence of autoimmune thyroid disease (AITD) leads to adverse reproductive outcomes and may severely affect pregnancy and offspring.

Screening programs

The controversy. Surprisingly, the need for routine thyroid evaluation in early pregnancy is far from being unanimously supported, possibly because these opinions are expressed by men. Since thyroid disease is common in the female population, checking thyroid function before or at the time of pregnancy diagnosis would appear to be essential. Certainly, thyroid testing is routine in fertility assessment and, because normal thyroid function is essential for normal intellectual development, screening for thyroid dysfunction is an important part of careful medical assessment.

Only recently has the American Association of Clinical Endocrinologists recommended thyroid function screening in all women seeking to become pregnant and/or during the first trimester of pregnancy. This screening should include determination of thyroid antibodies, which also represent important consequences for pregnancy outcome. The recommendation for aggressive case finding in women with increased risk for thyroid disease is nonsense in light of the high frequency of these disorders.

Choosing screening tests. Selecting only a single test for thyroid dysfunction screening in pregnancy may be inadequate. This is because of the controversy over the claim that serum TSH and T4 levels may not always be coincident. Half of so-called “hypothyroid” pregnant women display low T4 serum levels without high TSH and an equal number have high TSH serum concentrations with little change in serum T4. Indeed, very few women show both low T4 and high TSH levels. Usually, an abnormally high TSH concentration is of autoimmune origin and is associated with thyroid autoantibodies (TAb). The cause of the reported low T4 group remains speculative but may be related to iodine deficiency with T3 levels maintaining normal TSH or to unreliable testing assays. To avoid misinterpretation, it should be borne in mind that low TSH serum levels can also be a physiological response during the first trimester (see above).

While there is consensus on the utility of TSH determination, different opinions persist on whether serum TT4 or FT4 measurement should be the complementary test of choice (table 1). We prefer to use a T4 index with an appropriate binding assessment such as TBG or a T3 resin binding assay, since these indices give results in pregnancy similar to those in non-pregnant women.

Ultrasound. Although scintigraphy scans are contraindicated in pregnancy, routine ultrasound may be considered when nodular disease is suggested by history and examination. This procedure is useful not only for the diagnosis of thyroid neoplasms but also for the identification of maternal and fetal complications, such as goiter and fetal intrahepatic cholestasis of pregnancy.

TABLE 1. Routine recommendations for normal pregnancy

<table>
<thead>
<tr>
<th>Recommendation</th>
<th>Procedure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Screening</td>
<td>Recommended, especially when nodular disease is suggested by history and examination (250 µg/day; range, 200-500 µg/day)</td>
</tr>
<tr>
<td>Thyroid function</td>
<td>TSH and T4*: Interpretation should be trimester-specific</td>
</tr>
<tr>
<td>Presence of AITD</td>
<td>Anti-TPO and anti-Tg anti-thyroglobulin antibodies, anti-thyroid peroxidase antibodies, T4-thyron, TSH-thyrotropin.</td>
</tr>
<tr>
<td>Ultrasound</td>
<td>Recommended</td>
</tr>
</tbody>
</table>

AITD: autoimmune thyroid disease; anti-Tg: anti-thyroglobulin antibodies; anti-TPO: anti-thyroid peroxidase antibodies; T4: thyroxin; TSH: thyrotropin.

*See test: Once the diagnosis of pregnancy is confirmed, or preferably, before pregnancy occurs, isotope supplements should be started and screening for thyroid disease performed.
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only to characterize nodules and evaluate their growth characteristics but can also help establish a clinical diagnosis of Graves’ disease (GD) (by excluding nodules) or Hashimoto’s thyroiditis (based on typical heterogeneous patterning).

Diagnosis of suspected disease

Choice of tests. There is a similar controversy to that discussed above when choosing tests for the established pregnant patient under consideration. Once again, the most useful tests are determination of serum TSH, even though normal values are pregnancy-specific with an upper limit of < 2.5 µU/ml (fig. 1) and simultaneous TT4 (which is expected to be a maximum of 1.5 µg/dl above the normal range of non-pregnant women because of the increased TBG levels). A common practice in pregnancy has been determination of plasma FT4 to bypass the changes in TBG plasma levels. However, FT4 determination using commercial assays may be insensitive to the increase in serum transport proteins, which occurs during gestation, leading to false readings in the presence of high TBG. Moreover, there is no absolute value of FT4 that defines hypothyroxinaemia. In contrast, changes in TT4 during pregnancy are predictable and the assays do not depend on the problem of elevated TBG concentrations. A rougher higher reference range for TT4 during pregnancy can be calculated by multiplying the reference value of non-pregnant women by 1.5. Again, as mentioned earlier, in other centers, common practice is to use the relation between TBG and TT4 to calculate the free T4 fraction or T4 index (fig. 2).

Normal ranges. Interpretation of thyroid functions tests should be trimester-specific (fig. 1) although locally generated trimester-specific reference levels should be applied when available. Notably, the upper 95% confidence limit for serum TSH in the first trimester has been reported to be 2.5 µU/ml. TSH is known to decrease by 60-80% by week 10 and to recover slowly thereafter, but may not reach the normal range until gestation ends.

Diagnosis of hyperthyroidism

Diagnosis of hyperthyroidism may be supported biochemically when very low TSH serum levels are found (< 0.1 µU/ml) in the presence of concurrently elevated T4 concentrations. However, 10-20% of pregnant women may have low plasma TSH levels without concurrent thyrotoxic symptoms. Half of these women may detectable but subnormal serum TSH serum levels, whereas the other half has fully suppressed concentrations. When TSH is low, a trend in the T4 serum concentration can be helpful to differentiate transient physiological states from thyrotoxic conditions. Hence, more than just biochemistry is required for full confidence in the diagnosis. TSH receptor antibodies, eye disease, family history, goiter, weight loss, arrhythmias, and other factors need to contribute to the diagnosis.

Diagnosis of hypothyroidism

Evidence of an elevated serum TSH concentration makes the diagnosis of primary hyperthyroidism straightforward. The presence of thyroid antibodies is a useful confirmatory finding. According to some studies, serum TSH above 2.5 µU/ml in non-pregnant women should be used as a guide for thyroid dysfunction. Similarly, > 2.5 µU/ml is too high for the first trimester, and when serum TSH is > 4 µU/ml irrespective of the presence (or absence) of thyroid antibodies, there is no doubt about the presence of thyroid dysfunction. To confirm the diagnosis and severity, TT4 or FT4 must be measured. Normal pregnancy TT4 may be increased by 4-5 µg/dl (50-60 nmol/l) whereas FT4 should remain normal. Therefore, in early pregnancy any T4 levels below the normal range are suggestive of hypothyroidism, whereas in late pregnancy a FT4 values elevation of around 20-30% from pre-pregnancy values may be physiological.

AUTOIMMUNITY AND PREGNANCY

The immunology of pregnancy

Many changes in immune function develop during pregnancy, mostly initiated at the trophoblastic-uterine interface. For example, placental steroids and cytokines enhance the function of regulatory B cells and contribute directly to the modulation of immune reactivity in pregnancy. T cell function and antibody (Ab) secretion are both depressed during normal pregnancy, as reflected in the measurement of a wide variety of pathological antibodies and quiescence of many autoimmune diseases.

Thyroid autoantibodies

TAbs are present at birth in normal individuals but are suppressed by the normal immune system. However, almost 15% of the so-called normal population have easily detectable antibodies to thyroglobulin (Tg) and/or thyroid peroxidase (TPO), indicating that such suppression is faulty in a large number of people. Furthermore, post mortem studies have shown that the presence of serum thyroid antibodies is indicative of coincidental thyroiditis. Thyroid antibody levels fall significantly during pregnancy and then rebound in the postpartum once the suppression of pregnancy is lost.

Consequences of thyroid autoantibodies in pregnancy

The presence of TAbs in women of childbearing age has four main adverse consequences (table 2).

1. In euthyroid women, the presence of TAb has been correlated with early unexplained pregnancy
1. Loss of 30, probably related to a more generalized immune instability.

2. Women with TAbs before conception have an increased risk of developing hypothyroidism during pregnancy because of their reduced thyroid reserve consequent to their thyroiditis, which may negatively affect the development of the fetal CNS.

3. Irrespective of maternal thyroid function, the presence of TAbs is associated with higher rates of obstetric complications.

4. The presence of TAbs is directly related to the development of the postpartum thyroid syndromes.

The first reports of an association between an increase in early abortions and the presence of thyroid autoimmunity were published more than 15 years ago. The presence of TAbs in euthyroid women during the first trimester is now well known to be associated with a 2- to 4-fold increase in the abortion rate. A recent study found that pregnant women with positive anti-thyroglobulin Ab (anti-Tg), but without anti-thyroid peroxidase Ab (anti-TPO), also had an increased risk of very premature delivery in the absence of a significant association with TSH.

As mentioned earlier, there are several possible explanations for the association between TAbs and fertility-related adverse effects. Some authors have suggested that the presence of TAbs signals an autoimmune reproductive diathesis. TAbs may also indicate subtle thyroid dysfunction that negatively affects outcome.

TABLE 2. Effects of thyroid auto-antibodies (TAbs) in pregnancy

<table>
<thead>
<tr>
<th>Consequences of positive TAbs</th>
<th>Possible explanations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Delays in conception and consequently women are older when they conceive</td>
<td>Autoimmune reproductive diathesis</td>
</tr>
<tr>
<td>Early pregnancy loss</td>
<td>Generalized immune instability</td>
</tr>
<tr>
<td>Increased risk of developing hypothyroidism during pregnancy</td>
<td>Reduced thyroid reserve consequent to thyroiditis</td>
</tr>
<tr>
<td>Higher rates of obstetric complications</td>
<td>TAbs may be a marker of a generalized autoimmune imbalance. TAbs may also indicate subtle thyroid dysfunction that negatively affects outcome</td>
</tr>
<tr>
<td>Increased prevalence of postpartum thyroid syndromes</td>
<td>Predisposition to autoimmune diseases</td>
</tr>
</tbody>
</table>

Fig. 3. Thyroid autoantibody titer expressed as the mean at each time point of a study of a group of 33 euthyroid pregnant women who did not develop postpartum thyroiditis (group A) and a group of 33 euthyroid pregnant women who developed postpartum thyroiditis (group B). On average, Ab titers were two-fold higher in group B than in women who did not develop the disease (group A). The measurements were performed during each trimester and 3 and 6 months after delivery. Tg: thyroglobulin, TPO: thyroperoxidase. Modified from Stagnaro-Green et al.
of conception and women become older when they conceive\(^5\). It is also known that the higher women’s age, the greater the presence of TABs. These observations can be linked to the observed older age of women with poor obstetric outcomes. Lastly, women with TABs may have higher plasma TSH levels, reflecting subtle thyroid dysfunction. While some authors have found that levothyroxine (LT4) treatment did not prevent miscarriage in women with TABs\(^45\), others have reported prevention of pregnancy loss\(^46\). These data suggest that starting treatment with LT4 at the time of pregnancy may be appropriate to reduce the risk of miscarriage and prematurity in euthyroid anti-TPO positive women\(^47\).

The presence of anti-TPO in early gestation is also a useful marker for predicting the development of postpartum thyroid disease. Anti-TPO can usually be detected in 10% of euthyroid pregnant women at 14 weeks of gestation. After delivery, thyroid dysfunction (postpartum thyroiditis) occurs in 5-10% of all women. However, 30-50% of anti-TPO-positive women will develop postpartum thyroid dysfunction\(^48\). Maternal anti-TPO has also been related to intellectual impairment even when thyroid function was apparently normal, although once again thyroid deficiency may have been extremely subtle\(^49\). These data require confirmation.

**HYPERTHYROIDISM AND PREGNANCY**

**Epidemiology and etiology**

Hyperthyroidism during gestation is uncommon, secondary to the low fertility state, increased pregnancy loss, and the immunological changes that occur during pregnancy. The overall prevalence rate is about 0.1-0.4% of pregnant women, with GD accounting for 85-90% of all cases\(^50\). In addition, gestational transient thyrotoxicosis (GTT) occurs even more frequently but is not a disease\(^51\).

**Clinical features**

The presentation of thyroid hyperactivity during pregnancy may not be obvious because the symptoms may resemble gestational manifestations such as palpitations, excessive perspiration, dyspnea, and nervousness. Some signs, such as inadequate weight gain for gestational age, onycholysis, lid lag, muscle weakness and heart rate greater than 100 beats per minute (that does not decrease with the Valsalva maneuver), may help differentiate between thyrotoxicosis and the hypermetabolic state attributable to pregnancy\(^52\).

**Risks**

Hyperthyroidism affects pregnancy outcome, and serious complications may be associated. For the mother the risks include congestive heart failure, thyroid storm, preterm labour, pre-eclampsia and even death\(^53,54\), while the fetus may be stillborn, small-for-gestational age, or have congenital malformations\(^55\). Studies of pregnant women with the rare condition of thyroid hormone resistance syndrome demonstrated that their increased thyroid hormone levels, which are typical of this syndrome, were harmful to the normal fetus and caused a high miscarriage rate. From these studies, the potential for fetal loss in mild (subclinical) hyperthyroidism can be deduced. In addition, it can be inferred that expectant women may also be at risk if they have inadequately treated hyperthyroidism\(^56\).

Only isolated case reports of Graves’ associated ophthalmopathy and pregnancy have been published, since this complication rarely appears or worsens during pregnancy. In severe cases, surgical decompression may be warranted\(^57\).

**Gestational transient thyrotoxicosis**

GTT consists of biochemical hyperthyroidism in women with an otherwise normal pregnancy secondary to the thyrotrrophic effects of hCG. The prevalence of GTT depends on which ranges are used as normal for TSH and serum thyroid hormones but this disorder is clearly identified in approximately 2.3% of all pregnancies. In its most marked state, GTT is associated with morning nausea but improves spontaneously as pregnancy progresses. In the most severe form, high serum concentrations of hCG occur along with hyperemesis gravidarum –defined by severe nausea and vomiting leading to a 5% loss of body weight– dehydration and ketosis. Characteristically, whether symptomatic or not, GTT usually resolves spontaneously by 20 weeks’ gestation as hCG declines\(^58\). This disorder is, therefore, more frequent in multiple pregnancies, in which hCG levels tend to be higher. A tendency to reappear in subsequent pregnancies has also been observed\(^59\).

GTT must be differentiated from GD because the course, fetal risk, management and follow-up are different. GTT does not usually require specific treatment.

**Management**

Significant hyperthyroidism during pregnancy, whatever its cause (GD or nodules), must be treated. The most important issue to consider is that at least two patients are always involved\(^60\) (table 3).

**Antithyroid drugs.** In general, the treatment of choice in pregnancy is antithyroid drugs (ATD). Many physicians recommend the use of propylthiouracil (PTU) rather than methimazole (MMI) or carbimazole because of the widespread belief that this drug causes fewer side effects in the fetus. Very rare cases of aplasia cutis have been associated only with MMI and carbimazole\(^55,56\). However, if allergy or intolerance appears, it is still recommended that MMI be substituted. However, in Spain there are only two available ATDs, i.e., carbimazole or its metabolite MMI. Hence, PTU

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TABLE 3. Management of hyperthyroidism in pregnancy

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Indications</th>
<th>Recommendations</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antithyroid drug</td>
<td>Drug of choice: PTU (100-150 mg/8 h)</td>
<td>Monthly monitoring:</td>
<td>Maternal hormone target levels:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Clinical: mother and foetus.</td>
<td>TT4: 12.18 µg/dl or</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Ultrasound: foetus</td>
<td>FT4: 2.2-5 µg/dl.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Biochemical: tT4 (&lt;0.1 mU/L)</td>
<td>TSH: 0.1-0.4 µU/L (in late gestation)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>tSH and tT4*: mother</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Second trimester</td>
<td></td>
</tr>
<tr>
<td>Surgery</td>
<td>Uncontrolled maternal hyperthyroidism with high doses of ATD (over 300 mg of PTU or 40 mg/day MMI)</td>
<td>Should be temporarily used (no more than 4 weeks)</td>
<td>Side effects: increased risk of abortion or small-for-date infants</td>
</tr>
<tr>
<td>Beta-adrenergic blockers</td>
<td>Preparation for surgery</td>
<td>Should be used after week 32 if the disease is not controlled</td>
<td>Possible teratogenic effects</td>
</tr>
<tr>
<td>Radioactive iodine therapy</td>
<td>Contraindicated</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: See text.

Management of hyperthyroidism in pregnancy requires frequent monitoring. Treatment initially consists of anti-thyroid drugs. It is important to avoid hyperthyroidism in the fetus.

The initial dose of ATD depends on the severity of the disease. PTU is usually initiated at 100-150 mg/8 h guided by maternal T4 levels. The therapeutic aim is to maintain the “two patients” in a euthyroid state but fetal hypothyroidism must be avoided at all costs. ATDs pass through the placental barrier. Therefore, to avoid fetal hypothyroidism, the lowest dose possible to keep maternal T4 in the high normal range should be used. With this double objective in mind, once ATDs are started, “the patients” should be monitored every 4 weeks during gestation and the dose adjusted accordingly35. Monitoring consists of assessing maternal pulse, weight gain, thyroid size and measurements of TT4 (or FT4) and TSH with the recommended therapeutic target being TT4: 12-18 µg/dl (or FT4, 2-2.5 ng/dl)36. Normalizing TSH in early gestation to the non-pregnant range is not desirable, since the level in normal pregnancy is often low (fig. 1) and if significantly suppressed by excess thyroid hormone can remain suppressed for many weeks after regularization of peripheral hormones. The recommended range for TSH in patients on ATDs has been suggested to be 0.1-0.4 mU/L37. The median time to normalization of maternal T4 should be 7-8 weeks. Monitoring by TSH is more useful in late gestation when the disease is controlled.

The block and replacement approach (T4 plus ATD) should be avoided in pregnancy because of the difficulty of monitoring fetal thyroid function and the increased risk of producing goiter and hypothyroidism38.

When maternal hyperthyroidism is not controlled with high doses of ATD (over 300 mg of PTU or 40 mg/day MMI), surgery is often recommended in the second trimester. However, this happens rarely since the disease tends to improve as pregnancy progresses.

Managing the fetus. The management of the fetus requires serial ultrasound assessment for tachycardia, goiter, growth and hydromic changes. Fetal thyroid echo-Doppler may also be useful to evaluate gland function by a vascularization pattern. Goiter can usually be detected (when or if it is present) after week 32 but should be completely avoidable with appropriate disease management39.

Beta-adrenergic blockers. Beta-blockers should be avoided in pregnancy, or used temporarily for no more than 4 weeks until the ATD becomes effective in severe hyperthyrodisnym, or as preparation for surgery. Data in the literature have associated propranolol use in the first trimester with an increased risk of abortion and other effects such as small-for-date infants39.

Iodide. Information on use of large doses of iodides in pregnancy to control hyperthyroidism is scarce. Ancillary data consists of case reports. Low-dose iodide (6-40 mg/day of potassium iodide) has been used, leading to improvement in maternal thyroid function and normal neonatal outcome41,42. Because of the risk of fetal goiter, iodide use is not recommended except perhaps to prepare for surgery43.

Surgery. Subtotal or total thyroidectomy is indicated when, for any reason, ATDs fail to control the hyperthyroid disease44. The appropriate time is the 2nd trimester45. Ideally, surgery requires previous pharmacological treatment to normalize thyroid function but this may not be possible except with the use of beta-blockers and iodide.

Radioactive iodine therapy. During pregnancy 131I is contraindicated46 because of the possible teratogenic effects of radiation. However, the consequences of inadvertent administration of 5-10 mCi of 131I to pregnant women show that hypothyroidism occurs in only 3% of the fetuses46.

Special considerations in Graves’ disease

In general, the management of GD follows the same recommendations as those for any cause of hyperthyroidism.
Thyroid dysfunction in pregnancy

Management of Graves’ disease in pregnancy deserves particular consideration, bearing in mind the consequences of therapy on both mother and fetus. Stimulatory antibodies (TSHR-Ab) can stimulate both thyroids. The prevalence of neonatal hyperthyroidism (because of the passage of the antibodies) is not common (about 0.6–9.5% of cases) and can be predicted on the basis of high stimulating TSHR-Ab titers. In any event, neonatal GD is self-limiting and resolves in 5–10 months.

As previously stated, measurement of TSHR-Ab at 26 to 28 weeks’ gestation is recommended to evaluate the likelihood of neonatal hyperthyroidism. Fortunately, the prevalence of neonatal hyperthyroidism (because of the passage of the antibodies) is not common (about 0.6–9.5% of cases) and can be predicted on the basis of high stimulating TSHR-Ab titers. In any event, neonatal GD is self-limiting and resolves in 5–10 months.

As pregnancy progresses, the immune-privileged state disappears and a relapse, exacerbation or new onset of GD may occur between 4–12 months after delivery.

TABLE 4. Specific recommendations for Graves’ disease

<table>
<thead>
<tr>
<th>Condition</th>
<th>Recommendations for the mother</th>
<th>Recommendations for the fetus</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pregnancy in active GD</td>
<td>Monitoring TFT every month, and adjusting treatment accordingly. Assessment of TSHR-Ab titers (3rd trimester) to observe if the fetus is at risk.</td>
<td>Monitoring fetal pulse, which should not be tachycardic (&gt; 160 bpm). If maternal TSHR-Ab titers are positive, maternal thyroid function should be investigated after delivery.</td>
<td>Mother should continue with ATD</td>
</tr>
<tr>
<td>New diagnosis of GD during pregnancy</td>
<td>Medication should be restarted.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Relapse during early pregnancy</td>
<td>Maternal hyperthyroidism is impossible. Reassessment TSHR-Ab levels at the beginning to determine the probability of fetal or postnatal hyperthyroidism</td>
<td>If fetal tachycardia is detected in a fetus with maternal positive TSHR-Ab, initiating treatment with PTU 100-200 mg/8 h is recommended. As continuing LT4 supplementation to the mother to maintain maternal euthyroidism.</td>
<td></td>
</tr>
</tbody>
</table>

**Notes:**
- ATD: anti-thyroid drugs; PTU: propylthiouracil; T4: thyroxine.
- Management of Graves’ disease in pregnancy deserves particular consideration, bearing in mind the consequences of therapy on both mother and fetus. Stimulatory antibodies (TSHR-Ab) can stimulate both thyroids.
HYPOTHYROIDISM AND PREGNANCY

Epidemiology and etiology

Overall, the prevalence of hypothyroidism during pregnancy is approximately 2.5%, including both overt and mild (subclinical) cases. However, transiently high serum TSH levels in the first trimester can occur in two-thirds of women in certain populations, probably related to iodine deficiency. Notably, the percentage of pregnant women with increased serum TSH when they are TAb-positive is 40-60% compared with only 7-11% in matched non-pregnant Ab-positive women, revealing the stress placed on the thyroiditis-affected gland.

Determining the etiology of maternal hypothyroidism is important. The most common cause of hypothyroidism in women of reproductive age in the absence of iodine deficiency isAITD. A history of past total or subtotal thyroidectomy, radiiodine ablation or transient thyroiditis accounts for most of the remaining cases of hypothyroidism.

If maternal hypothyroidism is present, then the first trimester is the most critical time for fetal development, but if both maternal and fetal hypothyroidism occur (for instance in cases of TSHR blocking Ab or iodine deficiency) then all trimesters, and especially the third trimester, are critical periods. However, the presence of TSHR blocking Ab is a rare cause of maternal hypothyroidism (1 in 180,000 live births). These antibodies may be transferred to the fetus and cause intrauterine or transient neonatal hypothyroidism.

Clinical features

Hypothyroidism is associated with ovulatory dysfunction and consequently hypothyroid women have difficulty becoming pregnant. When hypothyroidism occurs, the signs and symptoms are similar to those in non-pregnant women, although only 20-30% of patients with overt hypothyroidism develop clear clinical features consistent with disease.

Risks

Hypothyroidism, even when mild, is classically associated with increased risks of anemia, gestational hypertension (pre-eclampsia or pregnancy-induced hypertension), fetal growth restriction, placental abruption, postpartum hemorrhage, cesarean section, perinatal mortality and neonatal morbidity. Perinatal delivery (before week 32) is three times more common in pregnant women with high TSH levels. Hypothyroidism is also associated with an increase in spontaneous abortions. Gestational hypertension occurs more often in overtly hypothyroid patients (36%) than in those with subclinical disease (25%) or in the general population (8%).

Danger to the future child

Another worrying aspect associated with maternal hypothyroidism (especially when present in early gestation) is the adverse consequence to fetal mental development. Several studies have shown clear evidence of the adverse effect of maternal hypothyroxinemia on neuropsychontellectual development in early childhood.

Management

Prevention of hypothyroid disease. Iodine supplementation is critical to prevent non-autoimmune maternal hypothyroxinemia during pregnancy, especially in iodine-deficient areas. Women of childbearing age with normally functioning glands should have an average iodine intake of 150 µg/day. During pregnancy and breast feeding, women should increase their daily iodine intake to 250 µg on average. LT4 therapy is required if, despite iodine supplementation, abnormal serum TSH levels are detected (table 1).

Levothyroxine supplements. As in non-pregnant situations, treatment of hypothyroidism depends on LT4 supplementation. To respond to the increased demands and to compensate the augmented binding capacity of thyroid hormone transport proteins, hypothyroid pregnant women already taking LT4 replacement therapy will require a dosage increase from 25% to 50% on average to maintain desirable TSH concentrations. Two-thirds of hypothyroid pregnant women need a dosage increase during the first trimester. In the second trimester there is usually a plateau in LT4 requirements but 25-40% of patients may require a further dosage increase in the third trimester.

The increase is at least partly dependent on the patient’s thyroid reserve. Hence, those patients with a history of a total thyroidectomy will be most dependent. Generally, patients with Hashimoto’s thyroiditis require a 25% increase in dosage while the increase should be 50% in full replacement therapy.

When high TSH is found for the first time during pregnancy, the test should be repeated but treatment...
Thyroid dysfunction in pregnancy

**TABLE 5. Management of hypothyroidism in pregnancy**

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Indication</th>
<th>Recommendations</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>LT4</td>
<td>High TSH</td>
<td>Dose: New diagnosis: start with 1.8-2 µg/kg (overt disease) or 100 µg/day (mild cases); i.e., TSH &lt; 10 mU/l</td>
<td>Maternal hormone target levels: FT4: 0.8-2.5 µg/dl; TSH: 0.5-2 µg/dl Possible drug interactions Postpartum period requires adjustments</td>
</tr>
<tr>
<td>LT4</td>
<td>Presence of TAbs*</td>
<td>Patients on LT4: increase dosage from 25% to 50% Monthly monitoring Clinical: mother Bioclinical (TSH and T4): mother</td>
<td></td>
</tr>
</tbody>
</table>

*FT4: free thyroxine; T4: thyroxine; TSH: thyroid stimulating hormone

**REFERENCES**

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