Prepregnancy Care and Patient Education Are Essential in Women with Thyroid Disease in Order to Prevent Pregnancy Complications

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SUMMARY

Background
Thyroid diseases are inconsistently reported to increase the risk for pregnancy complications. The objective of these authors was to study pregnancy complications associated with common and uncommon thyroid diseases.

Methods
Thyroid disease diagnosis (history of thyroid disease) and outcomes were derived from both electronic medical records and discharge summaries. Medication and laboratory data were not available. The authors analyzed singleton pregnancies (n = 223,512) from a retrospective U.S. cohort, the Consortium on Safe Labor (2002–2008). Multivariable logistic regression with generalized estimating equations estimated adjusted odds ratios (ORs) with 99% CIs. Main outcome measures included hypertensive diseases, diabetes, preterm birth, cesarean sections, inductions, and intensive care unit (ICU) admissions.

Results
Controls were pregnant women with no thyroid disease (n = 216,901). Primary hypothyroidism (n = 3183, 1.5% of pregnancies) was associated with increased odds of preeclampsia (OR, 1.47), superimposed preeclampsia (OR, 2.25), gestational diabetes (OR, 1.57), preterm birth (OR, 1.34), induction of labor (OR, 1.15), cesarean section (prelabor, OR, 1.31; after spontaneous labor, OR, 1.38), and ICU admission (OR, 2.08). Iatrogenic hypothyroidism due to thyroid surgery or ablation (n = 178, 0.1% of pregnancies) was associated with increased odds of placental abruption (OR, 2.89), breech presentation (OR, 2.09), and cesarean section after spontaneous labor (OR, 2.05). Hyperthyroidism (n = 417, 0.2% of pregnancies) was associated with increased odds of preeclampsia (OR, 1.78), superimposed preeclampsia (OR, 3.64), preterm birth (OR, 1.81), induction of labor (OR, 1.40), and ICU admission (OR, 3.70).

Conclusions
Thyroid diseases were associated with obstetric, labor, and delivery complications. Although the authors lacked information on treatment during pregnancy, these nationwide data suggest either that there is a need for better thyroid disease management during pregnancy or that there may be an intrinsic aspect of thyroid disease that leads to poor pregnancy outcomes.

ANALYSIS AND COMMENTARY

Thyroid diseases are not uncommon in women of reproductive age, with an estimation of 5% to 15% of women being affected, depending on geographic area, method of detection, and other, unknown factors. Euthyroid chronic thyroiditis is the most common thyroid condition, frequently suspected in the presence of a concomitant autoimmune disease, continued on next page
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a family history of thyroid disease, or palpation of a goiter and confirmed by the presence of serum TPOAb or TgAb. About 3% to 5% of patients with thyroiditis suffer from thyroid dysfunction, mainly subclinical hypothyroidism. In the majority of cases, this is detected during pregnancy if thyroid tests are ordered routinely. Universal screening for thyroid disease in pregnancy is controversial, but thyroid studies are strongly recommended at the time of the first obstetrical visit for women diagnosed with thyroid dysfunction before pregnancy, those on thyroid replacement therapy, and women with risk factors such as a family history of thyroid disease, autoimmune diseases, presence of goiter, and others (1,2). In the past two decades, numerous publications reported maternal, fetal, and neonatal complications and long-term neuropsychological deficits in children of mothers not properly diagnosed and treated during pregnancy. More alarming is the fact that 50% of women on thyroid replacement therapy have a serum TSH in the hypothyroid reference range when tested during pregnancy (3). The increase in demand for thyroid hormone in pregnancy is very well known, and due mainly to the stimulation of thyroid function by human chorionic gonadotropin (4). A recent paper showed that 80% of women with hypothyroidism who are on levothyroxine therapy before conception achieved a serum TSH within the reference range when tested during pregnancy (5). Encouraging reports recently published appeared to indicate that correction of maternal hypothyroidism in the second half of pregnancy achieved normal cognitive outcome in the offspring (6,7). The authors of the present study collected data limited to pregnancy outcome in a large multicenter study in women with a history of thyroid disease; however, no information on thyroid status was available at either conception or delivery. As recognized by the authors, the strengths of the study are its large size and comprehensive data collection from the hospital medical records from the intrapartum admission, allowing for the evaluation and adjustment for important confounding factors. In spite of the limitation in diagnostic and treatment data, the incidence and type of complications are similar to the ones reported in the literature.

Several studies in pregnant women with clinical hyperthyroidism and hypothyroidism showed that normalization of thyroid dysfunction has an impact in decreasing maternal and fetal complications. In this regard, the importance of prepregnancy education in our patients with thyroid disease is imperative; the endocrine community has educated women with prepregnancy diabetes about the importance of planning their pregnancies and achieving the best hemoglobin A1c values possible before conception, with a very successful result in reducing congenital malformations and other pregnancy complications. It is my opinion that the same approach should be used for women with thyroid disease, even if they are not contemplating pregnancy, keeping in mind that over 50% of pregnancies are unplanned. This educational approach should include women: (a) with active hyperthyroidism (contraception until euthyroidism is achieved); (b) on levothyroxine therapy, to achieve a serum TSH target close to 1 mIU/L before conception; (c) with euthyroid chronic thyroiditis, considering levothyroxine therapy if the preconception serum TSH is >1.5 mIU/L/ml; and (d) with risk factors for thyroid disease, offering a determination of thyroid tests (including both, TSH and TPOAb). In addition, these women should have thyroid-function tests at the time of pregnancy diagnosis and regularly throughout pregnancy and postpartum as clinically indicated. This approach should reduce pregnancy complications and prevent late offspring neurocognitive dysfunction.

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Prepregnancy Care and Patient Education Are Essential in Women with Thyroid Disease in Order to Prevent Pregnancy Complications

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