Familial Congenital Hypothyroidism Due to Inactivating Mutation of the Thyrotropin Receptor Causing Profound Hypoplasia of the Thyroid Gland

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Abstract

Thyroid gland agenesis is the most common cause of congenital hypothyroidism and is usually sporadic. We investigated a brother and sister from consanguineous parents, ascertained through systematic newborn screening, and initially diagnosed with thyroid agenesis. Careful cervical ultrasonography in both patients revealed a very hypoplastic thyroid gland. By direct sequencing of the thyrotropin receptor gene, we identified the substitution of threonine in position 553, in the fourth predicted transmembrane domain. The mutation was found homozygous in the affected siblings, and heterozygous in both parents and two unaffected siblings. Functional analysis in transfected COS-7 cells showed that it resulted in extremely low expression at the cell surface as compared with the wild-type receptor, in spite of an apparently normal intracellular synthesis. The small amount of mutated receptor expressed at the surface of transfected cells bound thyrotropin with normal affinity and responded in terms of cAMP production, but the in vivo significance of these data from overexpressed receptor in transfected cells is unclear. Of note, blood thyroglobulin was unexpectedly elevated in the patients at the time of diagnosis, a finding that might prove useful in refining etiologies of congenital hypothyroidism. (J. Clin. Invest. 1997, 99:3018–3024.) Key words: thyrotropin receptor • congenital hypothyroidism • thyroglobulin • thyroid agenesis

Introduction

The pituitary glycoprotein hormone thyrotropin (TSH) stimulates growth, differentiation, and function of the thyroid gland, and these effects are mimicked by cAMP agonists (1). TSH acts on the thyroid follicular cell through a specific receptor coupled to G proteins, whose hydropathy profile (2–5) displays a characteristic serpentine structure with seven transmembrane domains. Together with the other glycoprotein hormone receptors, LH/CG and FSH, the TSH receptor (TSHr) also presents, in its aminoterminal half, a large extracellular domain involved in hormone binding. The TSHr gene contains nine introns, with exon 10 coding for the entire serpentine structure (6). Besides stimulation of the cAMP cascade, the TSHr is capable of stimulating the phospholipase C–diacylglycerol–phosphatidylinositol pathway at higher concentrations of TSH, at least in man, and these effects are mediated by Gs and Gq proteins, respectively (1). In functional studies, the TSHr displays a significant level of spontaneous activity (i.e., stimulation of second messenger production in the absence of ligand) in experimental conditions where other G protein–coupled receptors are essentially silent (7–10). Mutations causing constitutive activation of the receptor have been found at the somatic level in hyperfunctioning thyroid adenomas (hot nodules) (7, 8), and at the germline level in familial autoimmune thyroid hyperplasia, a very rare condition (9). Reciprocally, inactivating mutations of the receptor have recently been described in patients with TSH resistance (11, 12) (see below).

Congenital hypothyroidism affects 1 in 4,000 newborns (13). Early thyroid hormone replacement therapy prevents the growth and development delay that develop postnatally in untreated patients, and systematic newborn blood screens for high TSH and/or low thyroxine (T4) have been implemented in most western countries. The majority of cases of congenital hypothyroidism are sporadic and are due to thyroid gland agenesis (athyrosis), or dysgenesis (ectopic and hypoplastic gland). These cases are attributed to a deficient embryogenesis of the thyroid; they do not show a clear tendency to recur within families of affected children and are believed to result from nongenetic causes, like environmental insults or stochastic events early in embryogenesis. A minority of congenital hypothyroidism cases, perhaps 10%, result from inborn errors of thyroid hormonogenesis, and show Mendelian inheritance, almost always of the autosomal recessive type (1). Homozygous or compound heterozygous mutations of the thyroglobulin (TG) gene, and of the thyroid peroxidase (TPO) gene, have been described in such families. In untreated patients, the increase in blood TSH activity often results in goiter, which may be present at birth (14).

It is expected that one subtype of inborn errors, if inactivating the TSHr, will not result in goiter, and instead resemble thyroid agenesis, except for showing autosomal recessive inheritance. A mouse strain called hyt/hyt presents with inherited autosomal recessive hypothyroidism and a hypoplastic gland (15). The affected mice were shown to harbor a homozygous mutation of the murine Tshr gene (16) causing defective TSH binding (17). In man, the concept of TSH unresponsiveness has been validated by clinical descriptions of patients (18, 19) and families (20) that fit the expected phenotype of low triiodothyronine and T4, and thyroid gland hypoplasia, in spite of...
high circulating levels of bioactive TSH. With the cloning of the human TSHr gene (3–5), molecular analysis of such cases became possible, but the search for mutations remained unsuccessful in previous studies (21). Recently, one patient with sporadic congenital hypothyroidism was shown to be a compound heterozygote for presumably inactivating mutations of the TSHr gene (22). On the other hand, compound heterozygosity for functionally tested, inactivating TSHr mutations was identified in siblings with resistance (i.e., partial unresponsiveness) to TSH without overt hypothyroidism (11, 12).

In this report, we describe a familial case of TSH unresponsiveness with congenital hypothyroidism due to a homozygous inactivating mutation causing very low expression of the TSHr at the cell surface, and profound thyroid hypoplasia.

Methods

Patients. Congenital hypothyroidism was identified through systematic screening on the fourth day of life in a boy whose blood spot TSH was > 130 mU/liter, and serum T4 was low at 1.2 μg/dl. The pregnancy had lasted 41 wk, and a cesarean section was performed because of transverse presentation. The birth weight was 4.30 kg, length 55 cm, and head circumference 41.7 cm. Physical examination and 99mTc scintigraphy failed to demonstrate the presence of thyroid tissue, and the patient was given a diagnosis of athyrosis. T4 therapy was initiated on the 9th d of life, and the neonatal course was otherwise uneventful. The family history (Fig. 1) revealed that the parents were second cousins, and that congenital hypothyroidism was previously diagnosed in a sister through an abnormal neonatal screen that came to our attention and for which normal values in our population (13). Serum TSH and free T4 were within normal limits in both parents and both unaffected brothers. Autoantibodies directed against TG, TPO/microsomal antigen, or the TSHr were not detected in the patients, nor in either parent or unaffected sibling. The parents consented to these studies. These data are summarized in Fig. 1.

Sequencing of the TSHr gene. DNA was extracted from peripheral blood using standard procedures. PCR amplification was designed to produce five overlapping fragments that cover the whole length of exon 10. Exons 1 through 9 were amplified individually using couples of intronic primers. The fragment encompassing the mutation identified in this family was generated using the following primers (positions 1567–1960): forward primer: 5′-tctggtatgccatcaccttc-3′; reverse primer: 5′-caggaaacagctatgacc-3′; forward primer: 5′-cagagacaccagcaaggtgag-3′; reverse primer: 5′-cagagacaccagcaaggtgag-3′; where the underlined sequences correspond to universal sequencing primers (−21M13 and M13RP1, respectively).

PCR was performed in a 100 μl final volume containing 1 μg DNA, 50 mM KCl, 10 mM Tris-HCl, pH 8.3, 2 mM MgCl2, 0.01% T4 was born. Two brothers and both parents were in good health, except for depression in the father that required lithium therapy. The family originated in Morocco and all children were born in Belgium.

Inconsistent with the diagnosis of athyrosis (23), serum TG was in the high normal range at the time of diagnosis in both patients (66 ng/ml in propositus, 84 ng/ml in affected sister). Careful ultrasonography in the propositus revealed a hypochogenic gland in normal position below the thyroid cartilage, whose transverse diameters were 5.2 and 4.9 mm for the left and right lobes, respectively, which corresponds to a very hypoplastic thyroid (24). Serum TG decreased over a 6-mo period while under T4 therapy. Treatment was interrupted for 6 wk in the sister at age 3.7 yr for investigative purposes. Physical examination showed no palpable thyroid, whereas serum-free T4 became very low, < 0.2 ng/dl. TSH increased to 186 mU/liter, and TG remained unchanged at 2.9 ng/ml. Like in her brother, ultrasonography showed a very hypoplastic gland in normal position, with transverse diameters of 5.6 and 4.6 mm for the left and right lobes, respectively.

Serum TG was found elevated at 53.7 ng/ml in the father, but was in the normal range at 17.1 ng/ml in the mother. TG levels in the unaffected brothers were 31.4 and 41 ng/ml, which is in the normal range for children in our population (13). Serum TSH and free T4 were within normal limits in both parents and both unaffected brothers. Autoantibodies directed against TG, TPO/microsomal antigen, or the TSHr were not detected in the patients, nor in either parent or unaffected sibling. The parents consented to these studies. These data are summarized in Fig. 1.

Figure 1. Thyroid blood tests and genotypes in the family. The parents are second cousins and the propositus is indicated by the arrow. Solid symbols indicate congenital hypothyroidism with apparently absent thyroid gland on physical examination and 99mTc isotopic scan. Blood tests were performed in the parents and four children. Values in unaffected siblings are at the time of diagnosis, and all measures are in the absence of thyroid hormone treatment. Normal values are given in brackets, except for serum TG, for which normal values in our population with borderline iodine intake are as follows (ng/ml): newborns, 2–106; 2 yr, 2–65; adults, 2–35. AutoAb, serum autoantibodies against TPO, TG, and the TSHr; ∼, below level of clinical significance; wt, wild-type TSHr sequence; M, Ala553→Thr (A553T) mutation as in Fig. 2.

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<th>1.80</th>
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<td>1.4</td>
<td>0.37</td>
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<td>TG, ng/ml</td>
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<td>17.1</td>
<td>31.4</td>
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gelatine, 0.2 mM dNTP, 5 U Taq Polymerase (GIBCO-BRL, Gaithersburg, MD), and 100 μM of each primer. The annealing temperature was 50°C. The PCR products were sequenced on both strands using the Thermosequenase fluorescent labeled primer cycle sequencing kit (RPN 2436; Amersham Corp., Arlington Heights, IL) according to the manufacturer’s recommendations. Samples were loaded on a 373 Stretch Sequencing Instrument (Applied Biosystems, Inc., Foster City, CA) and mutations identified using the Factory and Sequence Navigator Software (ABI Advanced Biotechnologies, Inc., Columbia, MD).

Cloning and expression of the mutated TSHR. A Ctn-BstII fragment (1604–2169) in the wild-type TSH receptor–pSVL construct was substituted by the mutant counterpart as previously described (7). Pfu enzyme was used to generate this fragment, in a 100 μl final volume containing 200 ng genomic DNA, 10 mM KCL, 20 mM Tris-HCl, pH 8.75, 10 mM (NH₄)₂SO₄, 2 mM MgSO₄, 0.1% Triton X-100, 0.1 mg/ml BSA, 0.2 mM dNTP, 5% DMSO, 2.5 U Pfu DNA Polymerase (Stratagene Inc., La Jolla, CA), and 0.5 μM of each primer. The annealing temperature was 52°C. The resulting constructs were verified by double stranded sequencing as described above.

COS-7 cells were grown in Dulbecco’s modified Eagle’s medium supplemented with 10% fetal bovine serum, 1 mM sodium pyruvate, 100 IU/ml penicillin, 100 μg/ml streptomycin, and 2.5 μg/ml fungizone, and seeded at a density of 150,000 or 300,000 cells/cm² dish. 1 d after transfection, triplicate dishes were used for flow immuno- cytofluorometry, cAMP determinations, and 125I-TSH binding studies; duplicate dishes were used for inositolphosphate determinations. Each experiment was repeated at least three times.

Binding and flow immunocytometry studies. 2 d after transfection, cells were washed twice with NaCl-free Hank’s buffer supplemented with 280 mM sucrose, 0.2% BSA, and 2.5% low fat milk, and were incubated at room temperature in fresh medium supplemented with 100,000 counts/ml 125I-TSH (35 μCi/μg, 40 U/mg TRAK assays; BRAHMS diagnostica, Berlin, Germany) and various concentrations of unlabeled TSH. 4 h later, the cells were rinsed twice with chilled Hank’s buffer and solubilized with 1 M NaOH. Bound radioactivity was determined in a gamma scintillation counter. Under these conditions, the radioactivity nonspecifically bound to COS-7 cells expressing the TSH receptor (defined as the radioactivity bound to the dish in the presence of 100 μg/ml unlabeled TSH) was identical to that bound to mock-transfected cells. In the absence of a consensus about the bioactivity of pure bovine TSH, we have expressed all TSH or TSH receptor concentrations in milliunits/milliliter, allowing for subtraction of ambient cAMP production by untransfected cells. C is thus the ratio between the cAMP level in cells expressing the receptor constructs over the cAMP level in the same number of cells transfected with the pSVL vector alone. C/R, normalizes C for the level of expression of each construct (see below). A measure of the relative constitutivity of the mutant receptor over the wild type is given by F = C mutant/C wild type. This value does not account for the differences in expression of the mutants at the cell surface. It gives a measure of relative constitutivity, at the actual level of expression achieved by individual receptor constructs. An estimation of specific constitutive activity (SCA) (i.e., constitutive activity normalized to equal level of expression) is given by SCA = F/R, where R is a measure of the expression of the mutant receptor relative to the wild type with R mutant/pSVL)/(WT-pSVL), the ratio between the fluorescence of cells transfected with the mutant and the wild-type receptor constructs, after subtraction of the background fluorescence displayed by mock transfected cells. The comparison of SCAs computed for each construct determines the relative constitutivity of molar amounts of the mutant over the wild type.

Statistics. Levels of significance were evaluated by a two-tailed unpaired Student’s t test, and P < 0.05 was considered significant.

Results

Familial mutation of the TSH receptor. Considering the clinical picture consistent with TSH unresponsiveness and the fa-
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As sole anomaly, we identified a GCC→ACC mutation (Fig. 2B) affecting the alanine residue at position 553, which was replaced by a threonine in the putative polypeptide chain (A553T). The Ala amino acid residue is highly conserved among the receptors for glycoprotein hormones of all mammalian species studied (2–5, 16), lending support to a pathogenic role of mutation A553T. Consistent with autosomal recessive transmission of the disease, the mutation was found homozygous in the propositus and his affected sister, whereas it was heterozygous in both parents. The two unaffected brothers were found to be heterozygous carriers of the mutation as well. The genotypes are mentioned in Fig. 1.

Expression of the mutated receptor. The functional characteristics of the mutated receptor were studied by transient expression in COS-7 cells. Cells transfected with a cDNA construct encoding the wild-type receptor or the empty pSVL vector were used as controls. Levels of expression at the cell surface were assessed by flow cytometry using a monoclonal antibody recognizing the linear epitope VFFEOQ (residues 354–359) (27, 29) of the extracellular aminoterminal domain. It demonstrated an extremely low level of expression for the mutant (Fig. 3). The comparison of saponin-permeabilized (Fig. 3B) and nonpermeabilized (Fig. 3A) cells showed that synthesis of the different constructs was quantitatively similar, which indicated that the mutant and wild-type receptors were equally recognized by the monoclonal antibody and that the low level

Figure 2. Familial mutation of the TSHr gene. B shows results from direct sequencing of PCR-amplified genomic DNA from peripheral blood in a normal unrelated control subject, in the propositus’ asymptomatic father, and in the propositus. It demonstrated a GCC→ACC mutation, affecting the alanine residue at position 553 (A553T), which was replaced by a threonine in the fourth putative transmembrane domain of the TSHr protein as schematized in A. Also represented in A is the location of the P556L mutation identified in the hyt/hyt mouse (see text). The genotypes of our patients and family are mentioned in Fig. 1.

Figure 3. Expression analysis of the A553T mutant by flow immuno-cytometry. Fluorescence intensity is expressed in arbitrary units, as a function of cell number (10,000 propidium iodide–gated cells) plotted on a logarithmic scale (see Methods). (A) Nonpermeabilized cells assayed after transfection with pSVL vector (pSVL), wild-type TSHr construct (wt TSHr), or A553T mutant TSHr construct (A553T). (B) Saponin-permeabilized cells transfected identically. Compared with pSVL-transfected cells, the shift in fluorescence of cells transfected with a receptor construct reflects the amount of receptor expressed at the cell surface (A) or present within the cell (B).
of expression at the cell surface resulted from a posttranslational event, perhaps a defect in the folding of the receptor resulting in abnormal routing to the plasma membrane. The expression of the mutant was identical when transfected cells were incubated at 30°C for 48 h before the functional assays (data not shown).

Binding studies were congruent with the FACS® analysis, and showed very low expression of TSH binding sites at the plasma membrane of cells transfected with the mutant construct, with an $R_t$ value of 0.03 mU/ml vs. 1.58 mU/ml for wild type in a representative experiment (Fig. 4). The observed dissociation constant of the mutant receptor was significantly smaller than wild type (0.21 mU/ml vs. 1.26 mU/ml, Fig. 4), which was confirmed in three independent experiments yielding values of 0.22±0.01 for $K_d$ and 0.018±0.01 mU/ml for $R_t$ (mean±SD). This may simply reflect the decrease in apparent $K_d$ that parallels the decrease in TSHr expression, as recently reported in stably transfected CHO cells (30), and as we observed in COS-7 cells (our unpublished observations). To exclude an additional effect of the mutation on the $K_a$, we performed binding studies on cells transfected with decreasing amounts of the wild-type receptor construct diluted in pSVL vector: when expressed at comparable levels, both constructs displayed comparable apparent affinities (Fig. 4).

**Activity of the mutated receptor on production of second messengers.** When overexpressed in COS-7 cells, the mutant receptor displayed a partial ability to stimulate cAMP accumulation in response to TSH (Fig. 5A). No systematic differences were observed as compared with the wild-type receptor for both $EC_{50}$ ($\sim 0.3$ mU/ml) and maximally active TSH concentration ($\sim 10$ mU/ml) (data not shown). In the absence of added agonist, the mutated receptor elicited a detectable level of cAMP accumulation that was smaller than in cells transfected with the wild-type receptor. As FACS® data allowed correction for transfection and expression efficiencies, we could compute the SCA as described in Methods. If the SCA of the wild type is arbitrarily put to 1, the A553T mutant displayed an SCA of $0.92±0.57$ (mean±SD) in five independent experiments. In spite of the dispersion of the values, which is essentially accounted for by the low expression of the mutated construct, we conclude that the mutant displays a constitutive cAMP stimulation activity similar to the wild-type receptor, in addition to its conserved cAMP response to TSH.

To confirm the recessive behavior of the mutation, we addressed the possibility of a dominant negative effect by cotransfection of the wild-type and the mutant constructs in various proportions, versus the wild-type construct mixed with pSVL vector to the same final amount of transfected DNA. For both the FACS® expression assay and the cAMP production assay, cotransfected cells behaved identically as the cells transfected with the wild-type receptor construct diluted in pSVL vector DNA (data not shown).

The mutated receptor was tested for its activity on the phosphatidylinositol–diacylglycerol regulatory pathway (Fig. 5B). In the absence of added agonist, we observed no significant stimulation of IP accumulation with the mutated as compared with the wild-type receptor. Stimulation by 100 mU/ml TSH elicited no response from the mutated construct as compared with the mock (pSVL)-transfected cells.

**Discussion**

Athyrosis is rarely, if ever, familial, and its cause has remained elusive in spite of its relatively high frequency. We show here that a genetic defect of the TSHr can cause a syndrome of apparent athyrosis that is transmitted as autosomal recessive.

The Ala$^{553}$→Thr missense mutation we identified in the TSHr gene affects a highly conserved amino acid residue, and cosegregates with the disease in this family. It is located in the fourth putative transmembrane domain of the receptor (Fig. 2A), and is reminiscent of the closely located Pro$^{556}$→Leu (PS56L) mutation of the murine TSHr, described in the hypothyroid hyp/hyp mouse, an animal model for TSH unresponsiveness (16). This mouse strain exhibits autosomal recessive congenital hypothyroidism with a normally located hypoplastic thyroid gland, as found in our patients. TSH unresponsiveness in these mice was found to result from defective TSH binding to a mutated receptor, which was apparently present at the plasma membrane (17). Unlike the mouse model however, TSH unresponsiveness in our patients is explained by the extremely low expression at the cell surface (Fig. 3A) of the A553T mutant receptor, in spite of synthesis in normal amounts (Fig. 3B).

Expression of TSH receptor constructs in COS-7 cells dif-

![Figure 4. Binding studies. $^125$I-TSH binding curves from a representative experiment with wild-type (wt TSHr) and mutated (A553T) TSHr constructs expressed in COS-7 cells (300,000 cells/dish) transfected with 500 ng construct DNA. Results are expressed as total number of receptors ($R_t$, mU/ml) and apparent dissociation constant ($K_d$, mU/ml). To compare wild-type and mutated constructs at a comparable level of expression, triplicate sets of dishes were also transfected with decreasing amounts of the wild-type construct diluted in pSVL vector DNA at 500 ng), yielding a low level of expression close to mutant, with a concomitant decrease in apparent $K_d$.](http://www.jci.org)
fers from the natural expression in thyrocytes in many aspects: (a) the level of expression is probably two to three orders of magnitude higher, as the consequence of transcription from a high number of copies containing a strong promoter; (b) the complement and concentrations of G proteins are likely to be different; (c) the characteristics of the membrane system and routing of membrane proteins are also different in a nonpolarized cell as opposed to the polarized thyrocyte. Despite these limitations, and perhaps because of some of them, overexpression of the A553T mutant in COS-7 cells provides pertinent information on structure/function relationships that add to the current molecular understanding of the TSHr, but whose in vivo relevance is unclear. The small number of receptors reaching the membrane seem to behave quite normally in terms of TSH binding and sensitivity to hormone for adenyl cyclase stimulation. The apparent decrease in $K_0$ is likely to result from the small number of receptors inserted correctly in the membrane (30), as this parameter normalizes when the wild-type and mutant receptors are expressed at similar levels (Fig. 4). The mechanism responsible for this effect is not understood, but it has also been observed for the PTH/PTHrP receptor (31). That the A553T receptor is competent for normal coupling to Gs is further indicated by its specific constitutive activity for cAMP generation, which is comparable with the wild type. Interestingly, the P556L mutation of the hyt/hyt mouse also displays constitutive activity. However, in this case, unresponsiveness to TSH must result from a structural defect since the receptor has been shown to reach the plasma membrane of transfected COS cells (17). As stated above, overexpression in COS-7 cells may nevertheless not reliably reflect the behavior of the TSHr in the thyrocyte, and the significance of the relatively robust residual cAMP production in COS-7 cells is unclear. If it participates in the observed phenotype, our findings may indicate that the blood levels of TSH sustained by our patients are limiting with regard to thyroid growth, as they remained comparable with the EC$_{50}$ of the mutant (0.3 mU/ml), which is much smaller than the pharmacological concentrations used on COS-7 cells (10 or 100 mU/ml) for maximal stimulation, and as human TSH is 5–10 times less potent on the human TSHr than the bovine TSH used in this study (32).

Although competent for coupling to Gs, the A553T mutant appears to have lost the capability to activate the phospholipase C–dependent regulatory pathway (Fig. 5 B). Dissociation of coupling to Gs and Gq has been observed previously in spontaneous TSHr mutants (8) and site-directed mutagenesis experiments (10). It agrees with a current model for G protein–coupled receptor activation, implying the existence of multiple discrete activated conformations.

The high TG in the patients at the time of diagnosis is surprising and not easily explained in the context of a very hypoplastic gland and congenital hypothyroidism (23). Apart from trivial causes of error and cross-reactivity with an unsuspected antigen in the TG assay, our patients show that significant expression of the TG gene can take place in vivo in spite of complete or near-complete TSH unresponsiveness. These findings may be confronted with the current knowledge of thyroid organogenesis and thyroid-specific gene expression. From the available evidence, it seems that development and migration of the embryonic thyroid in front of the trachea is independent of stimulation by TSH. Indeed, expression of thyroid-specific genes, including TG, TPO, and the TSHr, is subsequent to the observation of an in-place thyroid rudiment (33), which is believed to result from the interplay of at least three transcription factors, TTF1, TTF2, and Pax8 (34). However, it is tempting to suggest that normal TSHr function may be required to achieve a fully differentiated thyroid phenotype. The observation of cytological abnormalities in hyt/hyt mice, including overdistension of the endoplasmic reticulum, is compatible with such hypotheses (35). The disproportionately high plasma TG levels (considering the small size of the patients’ thyroids) could result from incomplete polarization of thyrocytes, abnormal routing of TG due to accumulation of misfolded mutant TSHr in the rough endoplasmic reticulum, or from intracellular leakage from dysplastic follicles. Testing of these hypotheses would require a biopsy of the thyroid from our patients, which we consider unethical.

The observed decrease and normalization of TG during T$_4$...
therapies are even more difficult to explain. It may simply correspond to the age-related decrease of serum TG observed in normal children from our population with borderline iodine intake (13). Alternatively, it is conceivable that irreversible full maturation of the thyroid could be normally dependent on auto-stimulation by thyroid hormones. Of note, TG levels did not increase when T₄ treatment was interrupted in the affected sister, in contrast with TSH that reached very high levels.

Unresponsiveness to TSH due to TSHr defects may represent a sizable subset of congenital hypothyroidism. It is probably rare even in the subgroup with familial occurrences of apparently absent thyroid tissue, as shown by the absence of any TSH mutations in a small series of patients with familial TSH unresponsiveness in previous studies (21). This leaves room for other, still unidentified genetic defects. The unexpectedly high serum TG levels in our patients suggest that measuring TG in all newborns diagnosed with hypothyroidism may help delineate further the etiologies of congenital thyroid malfunction in the future.

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References


