Coordination of hypothalamic and pituitary T3 production regulates TSH expression

Tatiana L. Fonseca,1 Mayrin Correa-Medina,1 Maira P.O. Campos,1 Gabor Wittmann,2 Joao P. Werneck-de-Castro,1,3 Rafael Arrojo e Drigo,1 Magda Mora-Garzon,1 Cintia Bagne Ueta,1 Alejandro Caicedo,1 Csaba Fekete,2,4 Balazs Gereben,4 Ronald M. Lechan,2 and Antonio C. Bianco1

1Division of Endocrinology, Diabetes and Metabolism, Miller School of Medicine, University of Miami, Miami, Florida, USA. 2Department of Medicine, Division of Endocrinology, Diabetes and Metabolism, Tufts Medical Center, Boston, Massachusetts, USA. 3Institute of Biophysics Carlos Chagas Filho and School of Physical Education and Sports, Federal University of Rio de Janeiro, Rio de Janeiro, Brazil. 4Department of Endocrine Neurobiology, Institute of Experimental Medicine, Hungarian Academy of Sciences, Budapest, Hungary.

Introduction

The negative feedback loop between circulating thyroid hormone and the neuroendocrine system (hypothalamus and pituitary gland) is the fundamental mechanism preserving thyroid hormone homeostasis. During iodine deficiency, there is decreased synthesis of thyroid hormone that leads to a reduction in plasma thyroxine (T4), rapidly activating secretion of hypothalamic thyrotropin-releasing hormone (TRH) and pituitary thyroid-stimulating hormone (TSH), which stimulates the thyroid to grow and normalize thyroid hormone synthesis and secretion (1). As with T4, a reduction in plasma 3,5,3′-triiodothyronine (T3), the biologically active thyroid hormone, also triggers the feedback mechanism and activates TRH and TSH secretion. This has been well documented in patients acutely given large amounts of propylthouracyl (PTU), which inhibits peripheral synthesis of T3 (2).

Given that the TSHB and TRH genes are negatively regulated by T3 (and not T4), it is crucial that T4 be converted to T3 in order to activate the negative feedback mechanism. The presence of type 2 deiodinase (D2), which converts T4 to T3 in the brain, is thought to be critical for this mechanism (3). D2 is coexpressed with TSH in pituitary cells (4) and is also expressed in astrocytes in thyroid hormone-sensitive regions of the brain, such as the cerebral cortex and hippocampus (5, 6), but its presence has also been documented in astrocytes in the mediobasal hypothalamus (MBH) (7). However, the highest expression of D2 in the MBH is in tanycytes, specialized ependymal cells lining the floor and infralateral wall of the third ventricle. In fact, tanycytes and TRH-expressing neurons in the paraventricular nucleus (PVN) are intermingled in the external zone of the median eminence (8, 9), providing the physical proximity for D2-mediated feedback signaling. Thus, hypothalamic D2 (astrocytic and tanycytic) has been proposed to have an important role in feedback regulation of thyroid hormone on hypophysiotropic TRH (10), but the relative importance of D2 expression in astrocytes versus tanycytes is not known.

Global inactivation of the D2 gene (also known as Dio2) in mice leads to elevated serum TSH and T4, but serum T3 remains normal, a phenotype compatible with an impaired transduction mechanism in T4-mediated TSH suppression (11). A very similar phenotype was also observed in mice treated with amiodarone, a noncompetitive inhibitor of D2 (12). However, neither the global D2 KO mouse nor the amiodarone-treated mouse models allow for a mechanistic understanding of the relative roles played by pituitary versus hypothalamic D2 in the thyroid hormone-mediated feedback mechanism. In addition, an unexpected finding was that both of these animal models exhibit normal PVN TRH expression, calling into question the role played by hypothalamic D2 in the T4-mediated feedback mechanism.

To address these questions, we developed the floxed D2 (D2f) mouse and used it here to generate 2 mouse strains: one exhibiting near-null D2 activity in the pituitary gland (referred to herein as pit-D2 KO mice), and one lacking D2 activity selectively in astrocytes (astro-D2 KO mice). We showed that the disruption in the T4-mediated mechanism of the pit-D2 KO mouse was bypassed by elevated serum T4 and reduced PVN TRH expression, defining a critical role for the MBH D2 pathway in the feedback mechanism. The phenotypic characterization of astro-D2 KO mice indicated that tanycytes, rather than astrocytes, are the cell type within the MBH, where this critical D2-mediated T4-to-T3 conversion takes place.
Results
Pituitary-specific D2 inactivation disrupts the hypothalamic-pituitary-thyroid axis (HPT). To determine the importance of pituitary versus hypothalamic D2 in HPT regulation, we used p-flox technology to allow for tissue-specific inactivation of the D2 gene based on selective expression of Cre recombinase (see Methods and Figure 1). We used mice expressing Cre recombinase under the pituitary-specific glycoprotein hormone α-subunit promoter (CGA-cre mice), which led to the generation of pit-D2 KO mice (CGA-cre/D2fl), a strain in which D2 activity was markedly reduced in the pituitary while remaining unaffected in the other D2-expressing tissues (i.e., cerebral cortex, hypothalamus, hippocampus, brown adipose tissue (BAT), and placenta; Figure 2, A–E, and Supplemental Figure 1M; supplemental material available online with this article; doi:10.1172/JCI61231DS1). D3 activity was unaffected throughout the brain (Supplemental Figure 1, A–C).

The pit-D2 KO mouse exhibited about 2.8-fold higher serum TSH levels and approximately 40% higher serum T4 levels compared with CGA-cre littermate controls, whereas serum T3 concentrations were not different (Figure 2, F, G, and I). No differences in 125I-T4 charcoal uptake were detected between pit-D2 KO and littermate control animals (Figure 2H), which indicates that the higher serum T4 was associated with elevated free T4 levels. However, that the pit-D2 KO mouse is systemically euthyroid was confirmed by studying tissue markers of thyroid hormone action. Total BW and individual organ weights in 2-month-old pit-D2 KO animals were similar to those of CGA-cre controls (Table 1). In addition, there were no differences in total skeleton bone mineral density (BMD), lean body mass (LBM), or oxygen consumption (VO2) (Tables 2 and 3), all sensitive markers of thyroid hormone action (13, 14). The very sensitive genetic marker of thyroid hormone signaling, liver D1 activity (15), was unaffected in pit-D2 KO mice (Figure 2J). The only noticeable difference was the approximately 20% lower body fat mass in pit-D2 KO animals (Table 2). Given that CGA is also expressed in gonadotrophs, we confirmed that the gonadal axis was pre-

Figure 1
D2 gene. The 2 loxP sites flanking the selenocysteine insertion site, base numbers, and amino acid positions are indicated. ATG, methionine; CDS, coding sequence.

Figure 2
Deiodinase activity and thyroid function tests in pit-D2 KO mice versus CGA-cre controls. (A–E) D2 activity in sonicates of pituitary, cortex, hypothalamus, hippocampus, and BAT (n = 3–5). (F and G) Serum TSH and T4 levels (n = 9–10). (H) 125I-T4 charcoal uptake in the serum (n = 9). (I) Serum T3 levels (n = 9–10). (J) Liver D1 activity (n = 3–5). *P < 0.05, **P < 0.001 vs. control.
served in pit-D2 KO mice, as shown by their normal serum levels of follicle-stimulating hormone (FSH) and luteinizing hormone (LH) as well as normal gonadal weight compared with controls (Supplemental 1, G–J).

We next hypothesized that reducing D2 expression in thyrotrophs would render these cells less sensitive to T4-mediated signaling, as has been proposed for the global D2 KO mouse (11). This was tested by assessing acute TSH responsiveness to T4 or T3. In control animals, administration of T4 elevated serum T4 approximately 7-fold and serum T3 approximately 3-fold, but reduced serum TSH by about 10-fold, to near-suppression levels (0.08 ± 0.002 ng/ml) (Figure 3, A–C). The drop in serum TSH occurred as a result of T4-to-T3 conversion in the thyrotroph and elevated serum T3. In contrast, administration of the same dose of T4 in pit-D2 KO animals only reduced serum TSH about 4-fold (0.5 ± 0.11 ng/ml), despite similar elevations in serum T3 and serum T4 compared with control animals (Figure 3, A–C). This can only be attributed to impaired thyrotrh T4-to-T3 conversion, given the identical T3-induced TSH suppression in both groups (Figure 3C).

Notably, there were no differences between groups in 30-minute TRH-stimulated TSH secretion: in all animals, serum TSH increased to similar levels, although the calculated fold response to TRH was greater in the control animals (~7 versus ~2-fold), given the higher baseline serum TSH in pit-D2 KO mice (Figure 3D). Furthermore, serum levels of T4 and T3 remained unaffected in these TRH-treated animals (Supplemental Figure 1, K and L).

To test D2-mediated T4 signaling in pit-D2 KO MBH, we studied PVN Trh mRNA levels in both mouse groups by in situ hybridization and found an approximate 50% reduction in pit-D2 KO mice (Figure 3E, F and F). However, Trh mRNA levels remained unaffected in the lateral hypothalamus of pit-D2 KO mice (Figure 3G). Given that MBH D2 remained unaffected in pit-D2 KO animals (Figure 2C), the elevated serum T4 and consequent increase in MBH D2–generated T3 explains the decreased TRH expression. This notion was supported by the finding that the difference in PVN Trh mRNA was minimized in control versus pit-D2 KO animals that were made systemically hypothyroid (105 ± 13 vs. 76 ± 20 AU; P > 0.05). Notably, the expression of key enzymes involved in processing the TRH peptide remained unaffected in the pit-D2 KO hypothalamus (Figure 3, H and I), which indicates that the decreased Trh mRNA is not neutralized by posttranslational mechanisms, thereby decreasing TRH peptide production. This finding illustrates a marked difference between the pit-D2 KO and global D2 KO mouse models, defining a major role for MBH D2 in the T4-mediated TRH feedback mechanism.

Reduced PVN TRH expression is expected to decrease hypothalamic stimulation of the thyrotroph; indeed, Tsha mRNA levels were reduced in pit-D2 KO mice (Figure 3J). However, pituitary Tshb mRNA levels were not affected in this mouse model (Figure 3K), presumably because the negative effects of T3 on TSHβ4 are greater than those on TSHα (16, 17). In other words, whereas Tshb was less stimulated by the hypothalamic, it was also less repressed by T3, due to inactivation of the local D2 pathway.

How could an elevation in serum TSH be reconciled with normal pituitary Tshb mRNA levels? We hypothesized that the reduction in pit-D2 KO PVN TRH expression prolonged TSH half-life and decreased its biological activity by altering the TSH glycosylation pattern (18). To test this hypothesis, we used a previously characterized in vitro assay to estimate pit-D2 KO TSH biological activity. Inducible cAMP production in CHO cells stably expressing TSH receptors approximately doubled with pit-D2 KO sera and decayed linearly with sera dilution (Figure 4, A and B). However, relative to the higher gravimetric amounts of TSH used in the assay (Figure 4C), pit-D2 KO TSH exhibited about 40% lower biological activity (Figure 4D). This is supported by our observations that thyroid structure, 2-hour 125I uptake, and mRNA levels of sodium/iode symporter (NIS) and thyroglobulin (Tg) were not different from those in control animals (Figure 4, E–I), despite the much higher serum TSH (Figure 2F). Notably, the pit-D2 KO thyroid gland was slightly enlarged, but the elevation in serum T4 and T3 observed 2 hours after bovine TSH administration was similar in both mouse groups (Figure 4, J and K).

Astrocyte-specific D2 inactivation does not affect HPT. The MBH contains high D2 activity levels, expressed in both astrocytes (7) and tanycytes (5, 6). To elucidate the contribution of astrocyte versus tanycyte D2 in the HPT, we created a second mouse strain expressing Cre recombinase driven by the glial fibrillary acidic protein (GFAP) promoter (GFAP-cre mice), a well-known marker for matured astrocytes that is rarely expressed in tanycytes (Figure 5A). Indeed, in the resulting astro-D2 KO mice (GFAP-cre/D2KO; see Methods), D2 mRNA levels in tanycytes were not altered compared with GFAP-cre littermate controls (Figure 5, B and C). However, astro-D2 KO mice exhibited a marked decrease in brain D2 activity in areas where D2 is expressed in glial cells (such as the cerebral cortex, hippocampus, and cerebellum), but not in the hypothalamus or pituitary gland, where D2 activity remained unaffected (Figure 6, A–E). In addition, D2 activity was also preserved outside the central nervous system (e.g., in BAT; Figure 6F). As with the pit-D2 KO mouse, D3 activity was not affected throughout the astro-D2 KO brain (Supplemental Figure 1, D–F).

Table 1

<table>
<thead>
<tr>
<th>Group</th>
<th>BW (g)</th>
<th>Brain</th>
<th>Epididymal fat</th>
<th>Liver</th>
<th>Muscle</th>
</tr>
</thead>
<tbody>
<tr>
<td>CGA-cre control</td>
<td>27 ± 3.8</td>
<td>14 ± 1.5</td>
<td>27 ± 10.2</td>
<td>44 ± 6.7</td>
<td>5.3 ± 0.7</td>
</tr>
<tr>
<td>Pit-D2 KO</td>
<td>29 ± 2.0</td>
<td>12 ± 0.58</td>
<td>24 ± 4.2</td>
<td>49 ± 3.1</td>
<td>5.6 ± 0.3</td>
</tr>
<tr>
<td>GFAP-cre control</td>
<td>28 ± 2.0</td>
<td>15 ± 1.8</td>
<td>13 ± 4.4</td>
<td>50 ± 6.3</td>
<td>5.4 ± 0.4</td>
</tr>
<tr>
<td>Astro-D2 KO</td>
<td>29 ± 3.4</td>
<td>15 ± 2.2</td>
<td>20 ± 1.4</td>
<td>45 ± 3.6</td>
<td>5.8 ± 0.3</td>
</tr>
</tbody>
</table>

Values are mean ± SD of 5 animals.

Table 2

<table>
<thead>
<tr>
<th>Group</th>
<th>BMD (mg/cm²)</th>
<th>LBM (g)</th>
<th>Fat mass (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CGA-cre control</td>
<td>51.8 ± 3.6</td>
<td>18.6 ± 2.4</td>
<td>6.3 ± 1.3</td>
</tr>
<tr>
<td>Pit-D2 KO</td>
<td>51.5 ± 2.4</td>
<td>19.0 ± 1.3</td>
<td>5.0 ± 0.7*</td>
</tr>
<tr>
<td>GFAP-cre control</td>
<td>55.7 ± 2.1</td>
<td>22.4 ± 1.8</td>
<td>7.2 ± 1.4</td>
</tr>
<tr>
<td>Astro-D2 KO</td>
<td>52.8 ± 1.4*</td>
<td>21.5 ± 1.8</td>
<td>6.2 ± 1.3</td>
</tr>
</tbody>
</table>

Values are mean ± SD of 5–7 animals. *P < 0.05 vs. respective control.
We found that astro-D2 KO mice had normal HPT and thyroid economy, as evidenced by their normal serum levels of TSH, T4, and T3; thyroid morphology; and PVN Trh mRNA expression (Figure 6, G–L). That the animals were systemically euthyroid was confirmed by analysis of multiple tissue markers of thyroid hormone action in 2-month-old animals, including BW, organ weights, LBM, fat mass, and VO2 (Tables 1–3). Finally, liver D1 activity was unaffected in astro-D2 KO mice (Figure 6M). In addition, we tested the relative sensitivity of TSH secretion to T4 or T3 and found that administration of either hormone suppressed serum TSH (Figure 6N), indicative of preserved HPT sensitivity to T4 in astro-D2 KO mice. As with pit-D2 KO animals, in both experiments, there were no differences between the serum levels of T4 and T3 achieved after administration of T4 or T3 (data not shown). An unexpected finding was a lower total body BMD in astro-D2 KO versus GFAP-cre control mice (Table 2), which suggests that brain D2 could be involved in bone metabolism or that D2 expression in certain GFAP-positive bone cells could play a previously unappreciated role in bone mass.

Discussion
The development and phenotypic characterization of 2 strains of tissue-specific D2 KO mice revealed a novel aspect of HPT regulation, namely, that a decrease in D2-generated T3 in the thyrotrophs can be compensated for by changes in hypophysiotropic Trh gene expression, presumably via an increase in D2-generated T3 in the MBH. Decreasing TRH expression in the pit-D2 KO mouse nullified any effect of selective pituitary D2 deficiency on Tshb gene expression, which remained unaffected (Figure 3K). However, the reduction in Trh expression (Figure 3, E and F) is likely to alter the pattern of TSH glycosylation, explaining the observed decrease in TSH biological activity (Figure 4, A–D).

MBH D2 expression is present in astrocytes and in tanycytes (5–7). Thus, our observation that thyroid economy was preserved in the astro-D2 KO mouse (Figure 6, G–I) indicates that tanycytic, and not astrocytic, D2 is required to transduce the...
T4-mediated negative feedback on PVN TRH expression. These mechanisms define a critical coordination between thyrotroph and tanycyte D2 pathways in the HPT and illustrate how preserving circulating serum T3 is indeed a centerpiece in thyroid hormone homeostasis.

The negative feedback mechanism in the thyrotroph is mediated by 2 important signals — serum T3 and serum T4 — via the D2 pathway (4). Thus, disruption of D2 in thyrotrophs would decrease their sensitivity to T4 and elevate serum TSH, which would in turn elevate serum T4 and T3 until a new steady state was reached at hyperthyroid levels. Alternatively, if serum T4 could bypass the thyrotroph and signal a negative feedback via the D2 pathway at the MBH, this would compensate for the lack of thyrotroph D2 and preserve euthyroidism. Our newly developed pit-D2 KO mice demonstrated that the latter mechanism is the case: serum T3 remained normal in these animals, and they were systemically euthyroid (Figure 2, I and J, and Tables 1–3). That the pit-D2 KO mouse exhibits increased T4 signaling in the MBH was confirmed by the reduction in PVN Trh mRNA (Figure 3, E and F). The observation that serum T4 was elevated in the pit-D2 KO mouse (Figure 3C) indicated that this MBH-based compensatory mechanism is less sensitive to serum T4, a phenomenon that could be explained by MBH D3 activity inactivating T4 en route to the tanycytes.

However, if the MBH D2 pathway compensates for thyrotroph D2 pathway inactivation, then why were elevated serum TSH levels observed in pit-D2 KO mice? The answer involves the posttranslational modification of the TSH molecule (i.e., TSH glycosylation), which is controlled by TRH (18). A reduction in TRH such as that seen during central hypothyroidism is expected to decrease TSH biological activity and prolong its half-life (18). In fact, pit-D2 KO TSH demonstrated approximately 40% decreased biological activity (Figure 4, A–D). This notion is strongly supported by the finding that most thyroidal parameters tested in the pit-D2 KO mouse were normal, including thyroid structure, 2-hour 125I uptake, and NIS and Tg mRNA levels (Figure 4, E and G–I), despite much higher serum TSH (Figure 2F). Thus, the new steady state reached by the pit-D2 KO mouse included normal pituitary-driven thyroidal activity, presumably because the elevation in serum TSH was mitigated by its reduced biological activity. Nevertheless, the pit-D2 KO mice exhibited goiter, with an approximately 60% increase in thyroid size (Figure 4F). This raises several possibilities, including selective TSH activation of thyroidal growth versus activity resulting from the expected alteration in TSH glycosylation, or earlier thyroid growth (before the new steady state was achieved).

Despite the elevation in serum T4, pit-D2 KO mice exhibited strong evidence for systemic euthyroidism, as documented by normal serum T3, and tissue euthyroidism, given the multiple T3-responsive parameters that were normal (Figure 2, I and J, and Tables 1–3). This indicates that in response to a disruption in pituitary T4 signaling, multiple adjustments take place in the HPT, eventually preserving systemic euthyroidism. In fact, the pit-D2 KO mouse model represented yet another example that the HPT is wired to maintain normal serum T3 levels, in addition to most animal models of isolated or combined deiodinase deficiency (19).
In the MBH, D2 has access to plasma-born or cerebral spinal fluid–born T4 to produce and release T3 that may then reach the PVN or the pituitary gland via portal blood (8). Previous studies in which MBH D2 activity is stimulated, either by LPS administration (20) or as a result of fatty acid amide hydrolase gene inactivation (21), indicate that D2-generated T3 can suppress TRH expression (22). In this region, D2 is reported to be expressed both in astrocytes (7), the only D2-expressing cell population that expressed GFAP-cre in the present studies (Figure 5A), and in tanycytes (5, 6). The preservation of MBH D2 activity in the astro-D2 KO mouse (Figure 6D) indicates that D2 is primarily expressed in tanycytes, although we cannot exclude with absolute certainty the possibility that selective ablation of D2 in hypothalamic astrocytes led to upregulation of D2 enzymatic activity in tanycytes to compensate for this loss. Nevertheless, by in situ hybridization histochemistry, no apparent increase in D2 mRNA was observed in astro-D2 KO versus GFAP-cre control tanycytes (Figure 5, B and C). We are confident that astrocyte D2 in the hypothalamus is markedly reduced in the astro-D2 KO mouse, given that D2 activity was dramatically reduced in every other location in which it is expressed in glial cells (including the cerebral cortex, hippocampus, and cerebellum), but remained unaffected in the pituitary gland and outside the central nervous system (Figure 6, A–C, E, and F). At the same time, we cannot exclude the possibility that MBH D2–generated T3 also reaches the thyrotrhop via portal blood and mediates, at least in part, the T4-mediated TSH negative feedback in pit-D2 KO mice.

TSH has been linked to significant extrathyroidal effects, namely, lipolysis (23, 24) and control of bone mass (25–27). Interestingly, the pit-D2 KO animals exhibited decreased fat mass and normal bone mass (Table 2), which could potentially shed some light on the roles played by TSH in these processes. However, the reduced TSH biological activity in the pit-D2 KO mouse prevents us from reaching definitive conclusions. At least it is clear that the elevated serum TSH we detected in the pit-D2 KO animals was not associated with increased bone mass, as would be expected by some (28, 29). Lastly, also unexpected was the observation that astro-D2 KO mice exhibited reduced BMD (Table 2). Although it is difficult to speculate on the underlying mechanisms (the global D2 KO has normal BMD; ref. 25), it is known that bone remodeling is under central control via the sympathetic nervous system (30, 31), a pathway that is affected by thyroid hormone and D2 expression (32, 33). Alternatively, D2 is expressed in bone and mouse osteoblasts (34) as well as in the chicken developing growth plate (35). Thus, given the GFAP expression in osteocytes and in cartilage chondrocytes (36), it is conceivable that selective D2 inactivation in these cells could have resulted in a phenotype that was not apparent in the global D2 KO mouse.

In conclusion, our characterization of the pit-D2 KO mouse revealed coordination between the D2-generated T3 pathways in the pituitary gland and MBH. In the absence of the pituitary D2 pathway, tanycyte D2 played a greater role within the MBH, reducing TRH expression and bringing the HPT to a new steady state in which serum T3 levels and euthyroidism were preserved. Included in this mechanism was an elevation in serum TSH and a reduction in its biological activity, which combined had a neutral net effect on thyroidal activity. Characterization of the astro-D2 KO mouse was instrumental in allowing us to define that tanycytic D2, not astrocytic D2, is a critical component of the feedback regulation mechanism of thyroid hormone on hypophysiotropic TRH.

Methods

Animals. To establish an in vivo mouse model for cell type–specific deletion of D2, first we generated D2loxP mice by inserting loxP sites (37, 38) into the D2 gene. The resulting D2loxP mice were then crossed with transgenic mice expressing Cre recombinase in a cell type–specific manner. The mouse D2 mRNA is unusually long (~6.5 kb) and contains a 8,071-bp-long single intron between codons 74 and 75 (39), making it risky to flox the whole exon 2 in view of the possible homologous recombination inside the floxed fragment. Therefore, the 5’ loxP site was placed into the 3’ region of the intron, while the 3’ loxP site was introduced to the 5’ region of the 3’ untranslated region (UTR) (Figure 1). According to this strategy, the loxP-flanked region contained the portion of the D2 coding region located in exon 2 (from codon 75 to the end of the protein) including the active center of the D2 enzyme required for catalytic activity (40). The presence of the 3’ loxP site in the 3’ UTR did not alter D2 activity (data not shown).

D2loxP mice were crossed with CGA-cre (B6SJL-Tg[Cga-Cre]3Sac/J; Jackson Laboratories; ref. 41) and GFAP-Cre (FVB-Tg[GFAP-cre]25Mes/J; Jackson Laboratories; ref. 42) transgenic mice, in order to eliminate the MBH D2 activity in the anterior pituitary (pit-D2 KO mice) and in GFAP-expressing astrocytes (astro-D2 KO mice), respectively. In all experiments, Cre littermates were used as controls. Except when stated otherwise, all mice were male, between 9 and 12 weeks of age at the time of the studies. Hypothyroidism was induced in some animals, as indicated, by adding 1% sodium perchlorate and 0.1% metimazol in drinking water for 30 days.

Body composition. Animals were fasted overnight, and total skeleton area, LBM, and fat mass were measured by dual energy X-ray absorptiometry (DEXA; Lunar Pixi). For the procedure, mice were anesthetized with ketamine/xylazine (200 mg/kg and 7–20 mg/kg) before imaging, as described previously (43).

Indirect calorimetry. Mice were individually housed and acclimatized in a calorimeter for 2 days before studies. Measurements of gas exchange were performed subsequently over the next 2 days as described previously (43). These studies were performed in a comprehensive lab animal monitoring system (CLAMS; Columbus Instruments), a computer-controlled open circuit calorimeter system. Metabolic profiles were generated based on real-time data obtained in successive 14-minute cycles. Studies were performed...
The sensors were calibrated against a standard gas mixture containing defined quantities of O₂ and CO₂ (Airgas). VO₂ was expressed as milliliters per kilogram per hour.

**TSH suppression studies.** TSH suppression studies were performed as previously described (11). Pit-D² KO mice and CGA-cre controls, or astro-D² KO mice and GFAP-cre controls, were placed on PTU (1 mg/liter drinking water) for 2 weeks. After blood was obtained from the orbital plexus (~150 μl), animals were injected s.c. with PTU (1 mg/100 g BW; 0.1 ml saline containing 0.01 N NaOH). 1 hour later, animals were injected s.c. with T₃ (1.2 μg/100 g BW), T₄ (3 μg/100 g BW), or vehicle. 4 hours later, new blood samples were collected, and animals were killed by CO₂ inhalation.

**TRH and TSH stimulation tests.** TRH or TSH tests were performed as previously described (44). Control and pit-D² KO mice received TRH (5.0 μg/kg i.p.; Sigma-Aldrich) or bovine TSH (250 mU/100 g i.p.; Sigma-Aldrich). Serum TSH, T₄, and T₃ levels were measured before and after TRH (30 minutes) or TSH (120 minutes) injection.

**Thyroidal ¹²⁵I uptake.** Thyroidal ¹²⁵I uptake was performed as previously described (19) after injection of Na¹²⁵I (15,000 cpm/g BW i.p.; NEN Life Science Products) into each animal. 2 hours later, mice were killed, and their thyroid glands were dissected and processed for radiometry. ¹²⁵I uptake was calculated as the ratio of thyroid cpm to total cpm and expressed as a percentage.
TSH biological activity. TSH biologically active was measured by bioassay using a line of CHO cells stably transduced with a human TSH receptor cDNA, as previously described (45). At the end of the incubation time, 220 μl of 10% trichloroacetic acid was added to each sample. cAMP was measured after adjusting to 6–7 pH, using cAMP 125I RIA kit (PerkinElmer).

Charcoal uptake. The thyroxine binding protein was determined using a modification of previously described methods (46). 10 μl mouse serum was diluted to 0.5 ml of PBS (pH 7.4) containing approximately 7,000 cpm 125I-T4, with specific activity of 135–165 μCi (5.00–6.11 MBq) per microgram (Perkin Elmer). Samples were allowed to equilibrate for 45 minutes at room temperature and then transferred to an ice water bath for 15 minutes. Prechilled 0.025% activated charcoal (0.5 ml; Sigma-Aldrich) solution in PBS was added, and samples were incubated on ice for an additional 15 minutes. Samples were spun in an Eppendorf centrifuge 5804R at 650 g for 15 minutes, and the charcoal-containing pellets were counted. Binding was calculated as the ratio of sample cpm to total cpm and expressed as a percentage.

Euthanasia and postmortem analyses. At the end of the experimental period, mice were euthanized by asphyxiation in a CO2 chamber. Brain, liver, skeletal muscle, and epididymal adipose tissue was dissected, weighed, and frozen in liquid nitrogen, and organ weights were expressed relative to total BW. The thyroid gland was dissected, weighed, and fixed in 4% paraformaldehyde for 24 hours at 4°C, and then embedded in paraffin, sectioned at 5 μm, and stained with H&E. Blood was collected, and serum levels of TSH, T4, T3, FSH, and LH were measured using a MILLIPLEX rat thyroid hormone panel kit (Millipore Corp.), according to the manufacturer’s instructions, and read on a BioPlex (BioRad).

In situ hybridization and analysis. In situ hybridization histochemistry was performed on every fourth section through the PVN or median eminence using a 741-base single-stranded [35S]UTP labeled cRNA probe for mouse TRH (47) or mouse D2 following previously described protocols (48). The coding region of the mouse D2 mRNA was amplified with Taq PCR (sense, ATGGGACTCCT-GAGCGTAGACTT; antisense, GCTAATCTAGAATTCATCTCTTGC) and cloned into a pGEM-T vector with its 5′ end at the T7 promoter. The sequence was confirmed by sequencing. The construct was linearized with NcoI and subjected to transcription with SP6 polymerase to generate the antisense strand. In vitro transcription was performed using SP6/T7 systems (Promega Corp.) and [35S]UTP (1250 Ci/μmole; PerkinElmer). The hybridization was performed under plastic coverslips in a buffer containing 50% formamide, a concentration of standard sodium citrate, 10% dextran sulfate, 0.5% sodium dodecyl sulfate, 250 ng/μl denatured salmon sperm DNA, and 5 × 106 cpm radiolabeled probe for 16 hours at 56°C. Slides were dipped into Kodak NTB autoradiography emulsion (Kodak) diluted 1:1 in distilled water, and the autoradiograms were developed after 3–7 days of exposure at 4°C. Autoradiograms were visualized under darkfield illumination using a COHU 4910 video camera (COHU Inc.). Images were captured with a color PCI frame grabber board (Scion Corp.) and analyzed with a Macintosh G4 computer using Scion Image. Background density points were removed by thresholding the image and integrated density values (density × area) of hybridized neurons on each side of the PVN measured in 3 consecutive sections of the midportion of the PVN where hypothalamic TRH neurons have been identified in mice (47).

Immunofluorescence for GFAP. 25-μm thick coronal sections from the hypothalamus cut on a cryostat were collected in PBS and treated with 0.5% Triton-X-100 plus 0.5% H2O2 in PBS for 15 minutes to increase antibody penetration. After rinses in PBS, sections were incubated in a mouse monoclonal antibody against GFAP (catalog no. MAB360; Millipore) overnight at a dilution of 1:2,000, diluted in 2% normal horse serum, 0.2% sodium azide, and 0.2% Photo-Flo. After rinses in PBS, sections were incubated in Cy3-conjugated donkey anti-mouse IgG (Jackson Immunoresearch) at 1:200 dilution for 2 hours. Sections were rinsed in PBS, mounted on glass slides, and coverslipped with Vectashield mounting medium (Vector).

Deiodinase assays. D1, D2, and D3 assays were performed in tissue sonicates in the presence of 10 mM DTT and 0.25 M sucrose as described previously (49). D1 and D2 assays were performed using 500 nM 125I-(5′) recombinant T3 (PerkinElmer) and 0.1 nM 125I-(5′) T4 (PerkinElmer), respectively, as substrates and 1 mM PTU for D2 measurements. D3 activity was measured as previously described (50) in the presence of 0.2 nM 125I-(5′) T3 (PerkinElmer) as substrate. Reactions were stopped by the addition of methanol, and the products of deiodination were quantified by UPLC (ACQUITY; Waters Corp.). Fractions were automatically processed through a Flow Scintillation Analyzer Radiomatic 610TR (PerkinElmer) for radiometry as described.

Gene expression analysis. Total RNA was extracted from pituitary and thyroid gland using RNAaquesious micro kit (Ambion; Life Technologies), according to the manufacturer’s instructions. The extracted RNA was quantified with a NanoDrop spectrophotometer, and 1.0 μg total RNA was reverse transcribed into cDNA using High Capacity cDNA Reverse Transcript kit (Applied Biosystems). Genes of interest were measured by RT-PCR (Bio-Rad iCycler iQ Real-Time PCR Detection System) using iQ SYBR Green Supermix (Bio-Rad) with the following conditions: 15 minutes at 94°C (hot start), 30–50 seconds at 94°C, 30–50 seconds at 55°C–60°C, and 45–60 seconds at 72°C for 40 cycles. A final extension at 72°C for 5 minutes was performed as well as the melting curve protocol to verify the specificity of the amplicon generation. Standard curves consisted of 4–5 points of serially diluted mixed experimental and control group cDNA. Cyclophilin A (CypA) was used as a housekeeping internal control gene. The coefficient of correlation was greater than 0.98 for all curves, and the amplification efficiency ranged between 80% and 110%. Results were expressed as the ratio of test mRNA to CypA mRNA. mRNA levels of the following genes were measured: Tsha, Tshb, Nis, Tg, peptidylglycine alpha-amidating monoxygenase (Pam), and glutaminyl-peptide cyclotransferase (Qpet).

Statistics. All data were analyzed using PRISM software (GraphPad). Unless otherwise indicated, data represent mean ± SEM. 1-way ANOVA followed by Student-Newman-Keuls test was used to compare differences among more than 2 groups; 2-tailed Student’s t test was used to compare differences between 2 groups. A P value less than 0.05 was considered significant.

Study approval. All experimental procedures were approved by the local IACUC (Miami, Florida, USA).

Acknowledgments

This work was supported by NIDDK grants R01 65055 and R01 37021, the National Science Foundation of Hungary (OTKA K81226), the Lendulet grant of Hungarian Academy of Sciences, and the European Community’s Seventh Framework Programme (FP7/2007-2013) under grant agreement no. 259772. We are grateful to Judith Molina for skilled technical assistance and Susy Bianco for assistance with cAMP assay.

Received for publication October 24, 2012, and accepted in revised form January 31, 2013.

Address correspondence to: Antonio C. Bianco, Division of Endocrinology, Diabetes and Metabolism, University of Miami Miller School of Medicine, 1580 NW 10th Avenue, Suite 602, Miami, Florida 33136, USA. Phone: 305.243.5631; Fax: 305.243.7268; E-mail: abianco@deiodinase.org. Or to: Ronald M. Lechan, Division of Endocrinology, Diabetes and Metabolism, Tufts Medical Center, 800 Washington Street, Box 268, Boston, Massachusetts 02111, USA. Phone: 617.636.8517; Fax: 617.636.4719; E-mail: rlechan@tuftsmedicalcenter.org.


