Thyrotoxicosis with absence of clinical features of acromegaly in a TSH- and GH-secreting, invasive pituitary macroadenoma


Department of Endocrinology, Diabetes and Metabolism, Cleveland Clinic Foundation, 9500 Euclid Avenue Desk F20, Cleveland, Ohio 44195, USA

1Pathology and Laboratory Medicine Institute 2Department of Neurosurgery and the Neurological Institute, Rose Ella Burkhartt Brain Tumor and Neuro-Oncology Center, Cleveland Clinic, Cleveland, Ohio 44195, USA

†R J Weil is now at Department of Neurosurgery, Geisinger Health System, 100 North Academy Avenue, Danville, Pennsylvania, USA

Correspondence should be addressed to P C Johnston
Email pcjohnston@doctors.org.uk

Summary

A 54-year-old woman presented with bi-temporal hemianopia, palpitations, and diaphoresis. An invasive pituitary macroadenoma was discovered. The patient had biochemical evidence of secondary hyperthyroidism and GH excess; however, she did not appear to be acromegalic. Surgical removal of the pituitary mass revealed a plurihormonal TSH/GH co-secreting pituitary adenoma. TSH-secreting adenomas can co-secrete other hormones including GH, prolactin, and gonadotropins; conversely, co-secretion of TSH from a pituitary adenoma in acromegaly is infrequent.

Learning points:

- This case highlights an unusual patient with a rare TSH/GH co-secreting pituitary adenoma with absence of the clinical features of acromegaly.
- Plurihormonality does not always translate into the clinical features of hormonal excess.
- There appears to be a clinical and immunohistochemical spectrum present in plurihormonal tumors.

Background

Thyrotropin (TSH)-secreting pituitary adenomas are rare and account for 1% or fewer of all pituitary adenomas. Approximately one-third of TSH-secreting pituitary adenomas co-secrete other hormones, the most common of which is growth hormone (GH) (16%) followed by prolactin (11%) and gonadotropins (1%). Conversely, in GH-producing pituitary adenomas, 13% have been shown to demonstrate immunopositivity to TSH (1) (2). Plurihormonal pituitary tumors are either morphologically monomorphous (single cells producing different hormones) or plurimorphous (different cells producing different hormones). Plurihormonal pituitary tumors seem to predict a higher risk of tumor recurrence, in comparison to tumors that secrete only one hormone; therefore, careful follow-up of this population is essential (3) (4). This report describes a rare case of a TSH/GH co-secreting pituitary adenoma with the absence of features of acromegaly.

Case presentation

A 54-year-old woman was noted on a routine visit to her ophthalmologist to have bi-temporal hemianopia.
She had reported difficulty with reading, but had not noted peripheral vision loss. Her medical history included atrial fibrillation, hypertension, cardiomyopathy, and sleep apnea. On review of symptoms, she reported hirsutism, feeling ‘hot’, palpitations, and diaphoresis. Physical examination revealed an obese woman with a BMI of 44, hirsutism, a moon-shaped face, and supraclavicular fullness. She had no goiter or dysthyroid eye disease. She had no overt features of acromegaly such as coarse facial features or broad fingers.

Investigation

Bi-temporal hemianopia was present, and magnetic resonance imaging (MRI) of the sella revealed a large pituitary macroadenoma measuring 2.3 cm and left cavernous sinus invasion, with suprasellar extension compressing the optic chiasm, principally to the left of midline. Initial laboratory testing revealed central hyperthyroidism: thyroxine (T₄) 11.4 (5.0–11.0 µg/dl), free thyroxine (fT₄) 2.1 (0.7–1.8 ng/dl), TSH 2.8 (0.4–5.5 µU/ml), and tri-iodothyronine (T₃) 185 (94–170 ng/dl), in addition to hypersomatotropism: insulin-like growth factor 1 (IGF1) 747 (87–267 ng/ml) and GH 1.58 ng/ml (Table 1).

<table>
<thead>
<tr>
<th>Test</th>
<th>Value</th>
<th>Normal range</th>
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<tbody>
<tr>
<td>fT₄</td>
<td>2.1</td>
<td>0.7–1.8 ng/dl</td>
</tr>
<tr>
<td>TSH</td>
<td>2.8</td>
<td>0.4–5.5 µU/ml</td>
</tr>
<tr>
<td>T₃</td>
<td>185</td>
<td>94–170 ng/dl</td>
</tr>
<tr>
<td>Prolactin</td>
<td>1.5</td>
<td>2.1–17.4 ng/ml</td>
</tr>
<tr>
<td>GH (basal)</td>
<td>1.58</td>
<td>0.01–0.97 ng/ml</td>
</tr>
<tr>
<td>IGF1</td>
<td>747</td>
<td>87–267 ng/ml</td>
</tr>
<tr>
<td>Cortisol a</td>
<td>6.6, 24.1, 15.8</td>
<td>&gt;18 µg/dl</td>
</tr>
<tr>
<td>ACTH (am)</td>
<td>16</td>
<td>8–42 pg/ml</td>
</tr>
<tr>
<td>24-h urine cortisol</td>
<td>16.3</td>
<td>&lt;45 µg/day</td>
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T₄, total thyroxine; fT₄, free thyroxine; TSH, thyroid-stimulating hormone; T₃, tri-iodothyronine; GH, growth hormone; IGF1, insulin-like growth factor 1; ACTH, adrenocorticotropic hormone.

*Short synacthen test: serum cortisol at baseline, +30, and +60 min respectively after administration of 250 µg cosyntropin.

Figure 1

Immunohistochemical staining results. (A) The adenoma shows diffuse staining for antibody against growth hormone (GH). (B) There is sparse and much weaker staining for antibody directed against TSH. Positively staining cells are brown.

Table 1  Pituitary hormonal profile at presentation.

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Outcome and follow-up

Three weeks later, she reported recrudescence of symptoms of hyperthyroidism (diaphoresis and palpitations), with recurrent evidence of biochemical hyperthyroidism and GH excess; methimazole and octreotide were re-introduced and 2 weeks later thyroid function normalized, T₄ 10.2 (5.0–11.0 µg/dl) and TSH 4.9 (0.4–5.5 µU/ml). Residual adenomatous tissue was present on sellar imaging...
identified in the suprasellar, intracranial space, displaced to the right by the tumor, and then identified passing through the diaphragm, to the normal gland within the sella; it was left undisturbed (Fig. 2B). Methimazole and octreotide were discontinued again on the day of surgery. Repeat immunohistochemistry of the resected tissue was identical to the initial histological findings. Ki-67 was 2.1 and < 5% of adenoma cells were p53 positive. Normalization of thyroid function and GH levels was observed post-operatively and has persisted, with complete resolution of the hyperthyroidism (Table 2). Post-operative MRI performed 6 weeks after surgery confirmed gross total resection of the intracranial component of the tumor; the infundibulum remained distorted to the right and is well visualized (Fig. 2C and D). At most recent follow-up, 3 years after the initial pituitary surgery, she has residual sleep apnea and continues to use ‘C-PAP’ nightly but is otherwise endocrinologically and neurologically normal.

Discussion

Plurihormonality does not always translate into the clinical features of hormonal excess. In addition to being distinct from one another with varying degrees of prominence, the clinical signs of hyperthyroidism and acromegaly can overlap. Patients with thyrotropic and somatotropic adenomas tend to present when the adenomas are large and frequently can be invasive and less amenable to surgical cure. Cure rates after surgery have been reported in around a third of patients with invasive tumors (5). Pre-operative somatostatin analog treatment can reduce tumor size and control thyroid and GH hypersecretion. Further options after surgery include radiotherapy and/or somatostatin analogs. Somatostatin analogs have been shown to result in euthyroidism in most cases of thyrotropinomas and ~50% achieve a reduction in pituitary mass size; dopamine agonists have also been used, but with more limited success in thyrotropinomas (6) (7).

In our case, the predominant symptoms were those of hyperthyroidism, although the clinical signs of hypertension, cardiomyopathy, and sleep apnea could be a clinical manifestation of acromegaly or morbid obesity as well. Interestingly, immunohistochemistry with antibodies directed against TSH showed rare and sparse positive cells. In contrast, although IGFI levels elevated to approximately three-times normal and showed diffuse robust staining against GH in the adenoma, the patient did not exhibit signs of acromegaly. One explanation is that the associated ‘classic’ signs often lag behind the
elevation of IGF1/GH. Another plausible explanation would be that even though the pituitary macroadenoma has been present for some time as indicated by its size (8), production of excessive somatotropins from the adenoma could have been a recent phenomenon. Secretion of GH with low bioactivity from a ‘silent’ somatotrope adenoma could also explain the relative absence of the clinical signs of acromegaly (9). Not all cases of biochemical and clinical TSH/GH co-secreting pituitary adenomas display immuno-positivity to TSH and/or GH. Negative immunostaining for TSH in co-secretory TSH/GH adenomas has been demonstrated, which could be due simply to low TSH levels in the adenoma (10). There appears to be a clinical and immunohistochemical spectrum present in plurihormonal tumors, and electron microscopy has been utilized by some to evaluate tumor cell lines in experimental animals (11). De novo transformation of a thyrotropinoma to a thyro-somatotrope adenoma appears to be exceedingly rare. The only case reported arose in a patient with hyperthyroidism from a thyrotropinoma, who developed evidence of GH co-secretion only after treatment with octreotide for 1 year. The authors speculated that a series of genetic mutations resulted in transformation of a monoclonal adenoma to a plurihormonal tumor (12). The PIT1 protein, which is a transcription factor whose presence is required in utero for normal development of the mammosomatotropes that produce GH and PRL, is also required for the maintenance of normal expression of GH, prolactin, and TSH after the separate embryonic origin of thyrotropins and has been postulated to play a role in the development of co-secretory pituitary adenomas (13). A recent study has demonstrated that the degree of β-TSH immunoreactivity appears to be associated with the degree of hyperthyroidism (14).

In summary, this case highlights an unusual patient with a rare TSH/GH co-secreting pituitary adenoma in whom symptomatic hyperthyroidism was the dominant and only clinical feature, without acromegaly. Immunohistochemistry was positive for GH but only rarely and sparsely positive for TSH. Secondly, while the patient had symptomatic relief of the signs and symptoms of hyperthyroidism for several weeks after what was thought to be gross total resection of a macroadenoma delimited by an elevated but anatomically intact diaphragm sella, the rapid recrudescence of hyperthyroidism led to early reappraisal and identification of residual component of tumor located wholly within the supra-diaphragmatic, intracranial space. Resection of the invasive, intracranial component of the adenoma led to resolution of biochemical evidence of acromegaly and the patient’s clinical hyperthyroidism.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement
All authors were involved in the preparation and writing of the manuscript.

References


