CASE REPORT

Relapse of hyperthyroidism after hemithyroidectomy in concurrent medullary thyroid cancer and Graves’ disease

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Abstract: We present a rare case of concurrent medullary thyroid cancer (MTC) and relapse of Graves’ disease (GD). A 26-year-old Chinese female suffered from GD for 1 year. Physical examination demonstrated mildly diffuse goiter with a palpable 1.0 cm nodule in the right thyroid lobe and exophthalmos. Before hemithyroidectomy, calcitonin was elevated and thyroglobulin was normal. After surgery, histopathological findings proved coexistence of MTC and GD. 16 months later, GD relapsed and radioactive iodine 131 therapy was given. Meanwhile, postoperative monitoring showed well controlled calcitonin and carcinoembryonic antigen. Our lesson demonstrates although concomitant of MTC and GD is very rare, yet after hemithyroidectomy for the single nodular MTC lesion, remnant thyroid lobe could be the source of GD relapse very soon. More importantly, the relapse of GD and the following radioiodine therapy and radiation exposure could have been prevented if a better therapy plan of total thyroidectomy was adopted in the first place, instead of hemithyroidectomy (Tab. 1, Fig. 2, Ref. 11).

Key words: medullary thyroid cancer, Graves’ disease, relapse, calcitonin, thyroidectomy.

The concurrency of thyroid cancer with Graves’ disease (GD) is very rare, occurring in 0.8–8.1 % of glands removed in the treatment of GD (1–4). Medullary thyroid cancer (MTC) is an uncommon and challenging malignancy, which accounts for only about 4 % of all thyroid cancer cases (1). We conducted a systematic literature search on PubMed database for articles in any language on the concomitancy of MTC and hyperthyroidism published from 1950 to 2010. We retrieved only 12 cases of concomitant MTC with hyperthyroidism (2–11), yet relapse of GD after surgical treatment of MTC has not been documented before. Here we report the 13th case, a 26-year-old Chinese female, who was the youngest and 1st such patient with GD relapse with in depth analysis of histopathology and immunohistochemistry.

Case report

This Chinese female patient was initially diagnosed with GD when she was 25 in another hospital. We could still retrieve her laboratory results at that time. She had elevated levels of triiodothyronine (T3, 648.0 ng/dL, reference 84.6–201.8), thyroxine (T4, 21.93 μg/dL, reference 5.13–14.06), free T3 (FT3, 27.90 pg/dL, reference 1.82–4.62) and free T4 (FT4, 7.91 ng/dL, reference 0.93–1.71), yet suppressed level of thyrotropin (0.01 μIU/mL, reference 0.27–4.20). She was given regular methimazole treatment for 1 year before she came to our hospital. Her past medical history was not significant. Physical examination demonstrated mildly diffuse goiter with a palpable 1.0 cm nodule in the right thyroid lobe and exophthalmus. Laboratory testing showed normal thyroid function tests. Thyrotrpin was 0.01 μIU/mL (reference 0.27–4.20). Postoperative imaging studies showed no evidence of residual disease. She remained euthyroid until 16 months after the operation when GD relapsed and radioactive iodine 131 therapy was given. Meanwhile, postoperative monitoring showed well controlled calcitonin and carcinoembryonic antigen.

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Fig. 1. A) 30 min after intravenous injection of technetium-99m pertechnetate, pre-surgical thyroid scan was performed, which demonstrated a cold nodule in the upper part of right thyroid lobe. B) 18 months after the operation when hyperthyroidism relapsed, a second thyroid scan was performed, which showed an enlarged left thyroid lobe with homogeneously increased uptake ability.
non-neoplastic parenchyma of the resected thyroid lobe exhibited typical GD features (Fig. 2B). By immunohistochemistry (Figs 2C–H), the tumor manifested positivity for Ct, chromogranin, synaptophysin and cytokeratin, and strong positivity for carcinoembryonic antigen (CEA), yet sporadic positivity for Ki-67. However, immunoreactivities for Tg and thyroid transcription factor-1 were negative. No lymph node metastases were detected. Because malignancy was revealed by pathology, a second surgery of complete thyroidectomy was discussed with the patient. Yet, the patient declined.

After hemithyroidectomy, she was given a regimen of 62.5 μg levo-thyroxine to obtain euthyroidism. For 15 months, her thyroid function was within normal ranges with Ct between 19.6 to 24.9 pg/mL and CEA between 0.2 to 1.2 pg/mL (reference 0.0–5.0). Ultrasound monitoring of the thyroid showed

<table>
<thead>
<tr>
<th>Author</th>
<th>Age (year)/Sex</th>
<th>Tumor size (cm) or TNM</th>
<th>Presurgical Ct (pg/mL)</th>
<th>Cause of hyperthyroidism</th>
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<tr>
<td>Pradeep (2)</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>Graves’ disease</td>
</tr>
<tr>
<td>Chao (3)</td>
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<td>NA</td>
<td>NA</td>
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<tr>
<td>Rieger (4)</td>
<td>61/M</td>
<td>T1 N0 M0</td>
<td>NA</td>
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<td>67/F</td>
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<td>NA</td>
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<td>Habra (5)</td>
<td>70/M</td>
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<tr>
<td>Brandle (6)</td>
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<td>Brandle (6)</td>
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<tr>
<td>Current report</td>
<td>26/F</td>
<td>1.4 cm</td>
<td>454 pg/mL</td>
<td>Graves’ disease</td>
</tr>
</tbody>
</table>

TNM: Tumor, lymph node and metastasis
NA: Not available

Fig. 2. A) Pathologically, hematoxylin and eosin stained section of the nodule demonstrated that the tumor cells with small, round hyperchromatic nuclei were moderately differentiated and were forming in shapes of nests or sheets. Morphologic features were suggestive to the diagnosis of MTC (magnification ×40). B) Hematoxylin and eosin stained other areas of the right thyroid lobe showed follicles lined by tall columnar epithelial cells and “scalloped” appearance on the edges of the colloid lumens, which were typical microscopic features of GD in histopathology (magnification ×40). C) Immunohistochemistry staining of the cancer lesion was positive for Ct (magnification ×40). D) Immunohistochemistry staining of the cancer lesion was positive for chromogranin (magnification ×40). E) Immunohistochemistry staining of the cancer lesion was positive for synaptophysin (magnification ×40). F) Immunohistochemistry staining of the cancer lesion was positive for cytokeratin (magnification ×40). G) Immunohistochemistry staining of the cancer lesion was strongly positive for CEA (magnification ×40). H) Immunohistochemistry staining of the cancer lesion was sporadic positive for Ki-67 (magnification ×200).

mally controlled FT3 (5.2 pmol/L, reference 3.5–6.5), FT4 (14.9 pmol/L, reference 11.5–23.5) and thyrotropin (3.69 μIU/mL, reference 0.20–5.00). Her preoperative calcitonin (Ct) was 453.8 pg/mL (reference <100.0), yet her thyroglobulin (Tg) was 9.14 ng/mL (reference <25.00). Ultrasound showed a 1.5×1.0×1.0 cm³ inhomogeneous nodule in the upper part of right thyroid lobe with left lobe of 1.8×2.1×5.1 cm³ and right lobe of 2.2×2.3×5.8 cm³, yet no enlarged lymph nodes were detected. Thyroid scan with technetium-99m pertechnetate (Fig. 1A) showed a cold nodule in the upper part of right thyroid lobe.

Right lobe hemithyroidectomy was performed, and the nodule was found to be a 1.4×1.0×0.8 cm³ gray-red lesion. Lymph node dissection of the right and central neck was also performed. Pathologically, the lesion demonstrated MTC cells forming in shapes of nests or sheets (Fig. 2A) with intact capsule. In addition, non-neoplastic parenchyma of the resected thyroid lobe exhibited typical GD features (Fig. 2B). By immunohistochemistry (Figs 2C–H), the tumor manifested positivity for Ct, chromogranin, synaptophysin and cytokeratin, and strong positivity for carcinoembryonic antigen (CEA), yet sporadic positivity for Ki-67. However, immunoreactivities for Tg and thyroid transcription factor-1 were negative. No lymph node metastases were detected. Because malignancy was revealed by pathology, a second surgery of complete thyroidectomy was discussed with the patient. Yet, the patient declined.

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tive findings. Nevertheless, 16 months later, nervousness, tremor, palpitation, weight loss despite increased appetite and increased sweating re-appeared. At that time, FT3, FT4 and thyrotropin were 22.0 pmol/L, 49.6 pmol/L and <0.01 μIU/mL, respectively. Levo-thyroxine was withdrawn immediately. During the following 2 months, her thyroid function decreased to some extent, yet still maintained at a high FT3 level of 10.4 pmol/L and FT4 level of 38.0 pmol/L, and a suppressed thyrotropin level of 0.01 μIU/mL. Ultrasound demonstrated an enlarged left thyroid lobe of 2.2×3.0×5.9 cm³ in size. 24-hour thyroid uptake rate of iodine was 56.5% (reference 10–40), and thyroid scan with technetium-99m pertechnetate (Fig. 1B) showed an enlarged left thyroid lobe with homogeneously increased uptake ability. At this time Ct, CEA and Tg were 17.9 pg/mL, 0.7 pg/mL and 2.3 ng/mL, respectively. She had a normal serum calcium, parathormone, and 24-hour urinary excretion of metanephrines, catecholamines and vanillylmandelic acid. GD relapse was diagnosed, and 6 mCi of radioactive iodine 131 (131I) was prescribed. After this therapy, euthyroidism was achieved within 3 months, and close follow-up was uneventful.

Discussion

MTC is a rare tumor derived from the parafollicular C cells of the thyroid. The simultaneous occurrence of thyroid cancer and hyperthyroidism varied from the published studies, while MTC was much less frequently found. We performed a systematic literature search on MEDLINE from 1950 to 2010 without language limitation. Key words were set as the combination of medullary thyroid cancer and Graves’ disease or toxic goiter or hyperthyroidism. Only 12 cases with such rare entities were retrieved (Tab. 1) (2–11). Our patient was the youngest, and also the 1st one who experienced GD relapse. Pre-surgically, we adopted serum Ct and Tg measurements as routine diagnostic evaluations of thyroid nodules. In the surgical management of the present case, hemithyroidectomy was foregone. Nevertheless, relapse of GD happened so quickly after merely 16 months, although no signs of MTC relapse or metastasis appeared. Therefore, the main lesson learned from this case is that the relapse of GD and the following radioiodine therapy and radiation exposure could have been prevented if a better therapy plan of total thyroidectomy was adopted in the first place, instead of hemithyroidectomy.

References


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