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Author(s)
Mussazhanova, Zhanna; Miura, Shiro; Stanojevic, Boban; Rougounovitch, Tatiana; Saenko, Vladimir; Shiraishi, Toshio; Kurashige, Tomomi; Shichijo, Kazuko; Kaneko, Kenichi; Takahashi, Haruo; Ito, Masahiro; Nakashima, Masahiro

Citation
Thyroid, 24(3), pp.593-598; 2014

Issue Date
2014-03-01

URL
http://hdl.handle.net/10069/34385

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Radiation-Associated Small Cell Neuroendocrine Carcinoma of the Thyroid: A Case Report with Molecular Analyses

Zhanna Mussazhanova,1,* Shiro Miura,2,* Boban Stanojevic,3,4 Tatiana Rougounovitch,3 Vladimir Saenko,3 Toshio Shiraishi,5 Tomomi Kurashige,2,6 Kazuko Shichijo,1 Kenichi Kaneko,7 Haruo Takahashi,7 Masahiro Ito,8 and Masahiro Nakashima1,2

Background: Neuroendocrine tumor (NET) of the thyroid other than medullary carcinoma is extremely rare. We describe here a case of calcitonin-negative small cell neuroendocrine carcinoma (SCNEC), which occurred in a thyroid gland that had previously been irradiated at high dose (60Gy) for pharyngeal cancer, with molecular analyses for follicular cell origin.

Patient Findings: The tumor cells were small with fine chromatin, inconspicuous nucleoli, and inapparent cytoplasm, and showed neuroendocrine architectures such as palisading, rosettes, and trabeculae. Mitotic figures were numerous exceeding 10 mitoses per 10 high-power fields. The tumor cells invaded into several vessels and metastasized to regional lymph nodes. Immunohistochemically, the tumor cells were strongly positive for neuroendocrine markers and thyroglobulin (Tg), a marker of thyroid follicular cells but negative for calcitonin and carcinoembryonic antigen (CEA). Expression of Tg and thyrotropin receptor (TSHR) were confirmed by quantitative real-time polymerase chain reaction (RT-PCR). Ki-67 labeling index was more than 70% in the tumor cells. Taken together, the tumor was diagnosed as SCNEC of the thyroid. Genetic analyses also revealed microsatellite abnormalities of the phosphatase and tensin homolog (PTEN) gene, suggesting that functional loss of PTEN contributes to carcinogenesis.

Conclusions: This is the first report describing a SCNEC of the thyroid with molecular analyses that provide evidence for a follicular epithelial origin.

Introduction

THYROID CANCER RISK after exposure to ionizing radiation is of public concern. Papillary thyroid carcinoma (PTC) is well known as the principal thyroid cancer cell type linked to radiation, especially after childhood exposure (1). The histological type of thyroid cancer associated with therapeutic external radiation exposure is most commonly PTC (2). Anaplastic transformation of PTC is also rarely induced by internal radiation with 131I after resection surgery for a primary tumor (3). We describe here a case of calcitonin-negative small cell neuroendocrine carcinoma (SCNEC) that occurred in a thyroid gland that had previously been irradiated at a high dose for pharyngeal cancer.

Neuroendocrine tumor (NET) of the thyroid other than medullary thyroid carcinoma (MTC), which is calcitonin positive, is extremely rare, and only one case of calcitonin-negative NET of the thyroid has previously been reported (4). That case was not associated with any history of radiation exposure and was histologically of the well-differentiated type, in which absence of mitotic activity, vascular invasion, and necrosis were noted. This is the first report describing a high-grade NET of the thyroid with molecular analyses with evidence of a thyroid follicular cell origin, and with alterations of the phosphatase and tensin homolog (PTEN) tumor suppressor gene.

Case Presentation

A 64-year-old Japanese male noticed a cervical mass. He was an ex-smoker of 30 cigarettes per day for 20 years. His family history did not include any endocrine disorders or any cancer syndromes. However, the patient had a past history of

*Z.M. and S.M. are jointly considered as first authors.
three primary cancers other than thyroid gland including esophageal cancer 18 years ago, squamous cell carcinoma of the lower pharynx 10 years ago, and moderately differentiated adenocarcinoma of the stomach 3 years ago. Surgical resections were performed for the esophageal and gastric cancers and no recurrence was evident so far. For the pharyngeal cancer, the patient underwent total resection of both the lower pharynx and larynx combined with left hemithyroidectomy and reconstruction by jejunal interposition. After the latter resection, he also received 60 Gy of external irradiation to the whole neck. One year before noticing his cervical mass, squamous cell carcinoma had recurred at his upper pharynx, which was treated by endoscopic mucosal resection, followed by 35 Gy of local external irradiation and chemotherapy with 5FU and CDDP.

Examination of the cervical mass by magnetic resonance imaging (MRI) disclosed a right thyroid mass and multiple swollen lymph nodes in the paratracheal and superior mediastinum, suggesting thyroid cancer (Fig. 1). Fine-needle aspiration was carried out, which showed malignant cells suspicious for undifferentiated carcinoma. Preoperatively, he was found to be profoundly hypothyroid: thyrotropin (TSH) 404.1 \( \mu \)IU/mL, free thyroxine (FT4) 0.13 ng/mL, free triiodothyronine (FT3) 0.30 pg/mL, and thyroglobulin (Tg) 215.9 ng/mL. The measurements of calcitonin and parathyroid hormone were not preoperatively performed. He subsequently underwent right thyroidectomy with dissection of the lymph nodes to remove the thyroid mass, which measured 5.4 cm \( \times \) 2.7 cm \( \times \) 1.9 cm, that occupied the right lobe of the thyroid gland (Fig. 1). He had no clinical evidence of tumor recurrence at the 1-year follow-up examination.

**Pathologic Findings**

The resected right thyroid gland was replaced by tumor cells and demonstrated few remaining nontumorous thyroid follicles. Tumor showed nested/nodular growth pattern with occasional foci of comedo-like necrosis, which was separated by fibrovascular stroma (Fig. 2). Neuroendocrine architectures, such as palisading, rosettes, and trabeculae, were clearly observed in the tumor nests (Fig. 2). The tumor cells were small in size (<3 lymphocytes) and oval to polygonal in shape with fine chromatin, inconspicuous nucleoli, and inapparent cytoplasms (Fig. 2). Mitotic figures were numerous exceeding 10 mitoses per 10 high-power fields (Fig. 2). Tumor cells invaded into several vessels and metastasized to regional lymph nodes. The pathologic tumor stage was pT3N1bM0.

**Immunohistochemical Profiles**

Immunohistochemistry was performed on formalin-fixed, paraffin-embedded (FFPE) tissue sections from the resected thyroid tumor. Antibodies used in this study are shown in Supplementary Table S1 (Supplementary Data are available online at www.liebertpub.com/thy). The immunohistochemical results are detailed in Table 1 and depicted in Figure 3, respectively. It was observed that the tumor cells were positive for neuroendocrine markers, such as synaptophysin and CD56, and weakly positive for chromogranin A. Further, the staining was strongly positive for thyroglobulin but negative for thyroid...
transcription factor-1 (TTF-1), both of which are markers of thyroid follicular cell. Immunoreactivity for calcitonin and carcinoembryonic antigen (CEA) was not observed in the tumor cells. The tumor was also weakly positive for p53, and Ki-67 labeling index was more than 70% in the tumor cells (Supplementary Fig. S1).

Molecular Analyses

Expression of TTF-1, Tg, and TSH receptor by quantitative real-time RT-PCR

Total RNA was extracted from FFPE tissues of the present case and two cases of PTC as well as their corresponding normal thyroid tissues, which acted as positive controls, using the High Pure RNA Paraffin Kit (Roche, Mannheim, Germany) according to the manufacturer's protocol. The cDNA was generated with the SuperScript III First Strand Synthesis SuperMix (Invitrogen, Carlsbad, CA) and analyzed by quantitative real-time polymerase chain reaction (RT-PCR) using Sybr Premix ExTaq qPCR primer assays (Takara Bio, Shiga, Japan) and the Thermal Cycler Dice Real Time multicolor PCR detection system (Takara, Shizuoka, Japan). The primers used in this study are summarized in Supplementary Table S2. Amplicons were designed to span introns to exclude genomic DNA contamination. PCR products were separated on 1% agarose gels and visualized by ethidium bromide staining (Fig. 4A). These assays demonstrated the expressions of both TSH receptor and Tg, but not of TTF-1, in the present case at the transcript level. All three genes were expressed by the two cases of PTC and/or their corresponding nontumor part of thyroid tissues.

Analysis for RET/PTC rearrangements and BRAF<sup>V600E</sup> mutations

Expression of RET/PTC1 and RET/PTC3 genes were assessed by RT-PCR as described in a previous report (5). Neither RET/PTC1 nor RET/PTC3 was demonstrated in the present case. Furthermore, tumor DNA was amplified to generate products covering the common mutualational hotspots of BRAF, including a portion of exon 15 and and sequenced on an ABI PRISM 3130 automated capillary DNA Sequencer (Applied Biosystems) using the BigDye Terminator Cycle Sequencing Ready Reaction Kit version 3.1 (Applied Biosystems). No BRAF mutations were detected in the present case.

Loss of heterozygosity analysis of the phosphatase and PTEN tumor suppressor gene

Using a Tissue Mini DNA Purification Kit (Epicentre Biotechnologies), genomic DNA was extracted from FFPE tissues.
of both the tumor and its corresponding lymph node without metastasis from the present case as well as a case of MTC and a case of colonic NEC, which were used for loss of heterozygosity (LOH) analysis by PCR. PCR reactions were routinely performed in a final volume of 25 μL containing 10 ng of sample DNA with the hot start ExTaq polymerase (Takara). Five microsatellite markers were used to determine LOH in the 10q23 region and its flanking polymorphic genetic markers as described in a previous report (6). PCR was performed under the following conditions: initial denaturation at 95°C for 5 minutes followed by 37 cycles of amplification (94°C for 30 seconds, 56°C for 30 seconds, and 72°C for 30 seconds), and a final extension at 72°C for 10 minutes, followed by a 4°C cooling step. Aliquots of the PCR products were electrophoresed on 12% denaturing polyacrylamide gels at 50V for 24 hours. After electrophoresis, the allelic band intensity was detected using a commercially available ethidium bromide solution, and was captured and analyzed using a BioDoc-It Imaging System (UVP, Inc., Upland, CA). LOH was scored when the intensity of one allele was absent or at least 75% reduced in the tumor alleles compared to the corresponding wild-type alleles in the normal tissue.

The present case showed microsatellite abnormalities in three of the five markers, including a biallelic loss at D10S541 and two monoallelic losses at D10S2491 and D10S583, while neither thyroid medullary carcinoma (MC) nor colonic neuroendocrine carcinoma (NEC) showed any microsatellite abnormalities in these markers.

**Discussion**

This article describes a case of SCNEC of the thyroid that occurred in a patient who had a past history of radiation therapy for hypopharyngeal cancer. In the thyroid gland, although MTC is well known as a NET arising from the calcitonin-producing parafollicular cells (C-cells) that accounts for up to 10% of all thyroid cancers, only one case of calcitonin-negative
nonmedullary NET has previously been reported, which expressed Tg, synaptophysin, and chromogranin A, but not calcitonin by immunostaining, indicating a follicular cell origin (4). In the present case, the expression of three markers of neuroendocrine differentiation, but not of calcitonin nor CEA, were demonstrated. We also confirmed the expression of thyrotopin receptor (TSHR) and Tg at the transcription level, providing further evidence for a follicular cell origin of this tumor.

The expression of TTF-1 was not detected in the present case at both mRNA and protein levels. TTF-1 expression is highly restricted to epithelial tumors arising in the thyroid and lung. Among pulmonary NETs, it has been documented that TTF-1 expression is found in 35–50% of typical carcinoids, 100% of atypical carcinoids, 50% to 75% of large cell neuroendocrine carcinomas, and 81% to 95% of small cell carcinomas (7–9). Among thyroid NETs, MTCs seem to consistently express TTF-1 (7,10), which is plausible because this protein is expressed in C-cells (11,12). TTF-1 is a homeodomain-containing transcriptional factor that binds to and activates the promoters of thyroid-specific genes including TSHR, Tg, and thyroid peroxidase genes in follicular cells (13). Therefore, during carcinogenesis of SCNEC of the thyroid, the TTF-1 gene may be truncated, and the expression of some thyroid-specific genes, such as TSHR and Tg, may be regulated in a TTF-1 independent manner.

NETs may arise in most organs of the body and share many common pathologic features. The 2004 World Health Organization (WHO) classification recognizes four major types of lung NETs: low grade/typical carcinoid, intermediate grade/atypical carcinoid, high grade/large cell neuroendocrine carcinoma, and small cell lung cancer (14). More recently, according to the 2010 WHO classification scheme, gastroenteropancreatic (GEP)-NETs are separated into well-differentiated NETs and poorly differentiated NECs of the small or large cell type (15). They are also divided according to a grading system, which is based on the Ki67 proliferation index or mitotic count. According to these criteria, the present case is a high-grade (G3) NEC of small cell type. To the best of our knowledge, this is the first report of calcitonin-negative high-grade NEC of the thyroid.

Acknowledgments

This work was supported in part through Nagasaki University Global Center of Excellence (COE) program “Global Strategic Center for Radiation Health Risk Control” and by Grant-in-Aid for Scientific Research from the Japanese Ministry of Education, Science, Sports and Culture (No. 18590334).

Author Disclosure Statement

The authors of this manuscript have no competing financial interests. They have no commercial associations that might create a conflict of interest in connection with the submitted manuscript.

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Address correspondence to: Masahiro Nakashima, MD, PhD Department of Tumor and Diagnostic Pathology Atomic Bomb Disease Institute Nagasaki University 1-12-4 Sakamoto Nagasaki 852-8523 Japan
E-mail: moemoe@nagasaki-u.ac.jp