# Four cases of medullary thyroid carcinomas associated with multiple endocrine neoplasia 2B with rearranged during transfection codon M918T mutation in the same family

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Abstract. Multiple endocrine neoplasia (MEN) with medullary thyroid carcinoma (MTC) is associated with rearranged during transfection (RET) mutations. The authors encountered four cases of MTC-related MEN type 2B (MEN2B) with RET codon M918T mutation in one family. Case 1 included a 19 year-old male diagnosed with MTC with lung metastases. Genetic testing revealed an RET codon M918T mutation, which indicated MEN2B. The patient responded partially to vandetanib and the disease has shown no progression in 25 months. Case 2 involved the mother of the patient in Case 1. She underwent total thyroidectomy (TT) when diagnosed with MTC-related MEN2B at 12 years of age, but was not counseled adequately. Cases 3 and 4 involved the sisters of the Case 1 patient and were assessed after Case 1 was diagnosed. Genetic testing revealed the same mutation. Case 3 was diagnosed with MTC and underwent TT. Case 4 was asymptomatic but underwent prophylactic TT; histopathologic examination revealed MTC tissue. Prophylactic TT prevented MTC from being detected at an advanced state. Genetic counseling is essential in treating MEN2B. The mother was uninformed about the genetic characteristics of MEN2B, delaying the detection of

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*Abbreviations:* MTC, medullary thyroid carcinoma; MEN, multiple endocrine neoplasia; RET, rearranged during transfection; TT, total thyroidectomy; CEA, carcinoembryonic antigen; Ctn, calcitonin; CT, computed tomography; PHEO, pheochromocytoma

*Key words:* medullary thyroid carcinoma, multiple endocrine neoplasia, RET codon M918T mutation, genetic counseling, prophylactic total thyroidectomy

MTC in her children. The present study reaffirms the importance of family history and screening.

#### Introduction

Medullary thyroid carcinoma (MTC), which accounts for 1-2% of thyroid carcinomas, occurs sporadically or in a hereditary form associated with two types of multiple endocrine neoplasia (MEN), MEN2A and MEN2B, as well as familial MTC (FMTC) (1-7). MEN is inherited in an autosomal dominant manner (4,5). The rearranged during transfection (RET) germline mutation is associated with MEN2A, MEN2B and FTMC and most patients with MEN2B have RET codon M918T mutation (2-7). The RET codon M918T mutation is the highest risk factor for MTC, according to American Thyroid Association (ATA) guidelines (8). Although the diagnosis of MEN2B at an early age is optimal, the majority of patients are diagnosed after reaching an advanced state (9). MTC and MEN2B patients as well as their relatives must be examined carefully due to the hereditary characteristics of this disease. Genetic counseling is necessary in numerous cases. The authors of the present study encountered four cases of MTC with MEN2B in the same family. Sequencing by synthesis was adopted for genetic testing of all four cases (10).

# **Case reports**

*Case 1*. The first case was that of a 19-year-old male presenting with cervical swelling, dyspnea and chronic diarrhea. Peculiar physical features included neuromas of the tongue (Fig. 1A) and eyelid (Fig. 1B), as well as marfanoid habitus (Fig. 1C). A blood test revealed high levels of carcinoembryonic antigen (CEA) and calcitonin (Ctn). Computed tomography (CT) showed unresectable tumors of the thyroid, bilateral neck lymphadenopathy (Fig. 2A) and multiple nodules in both lungs (Fig. 2B). The trachea was narrowed and deviated due to a thyroid mass (Fig. 2C). Histologic examination confirmed a diagnosis of MTC (cT4bN1bM1, stage IV). Due to lung metastases, vandetanib, a tyrosine kinase inhibitor, was administered. The daily dose of vandetanib was 300 mg. Cardiovascular monitoring

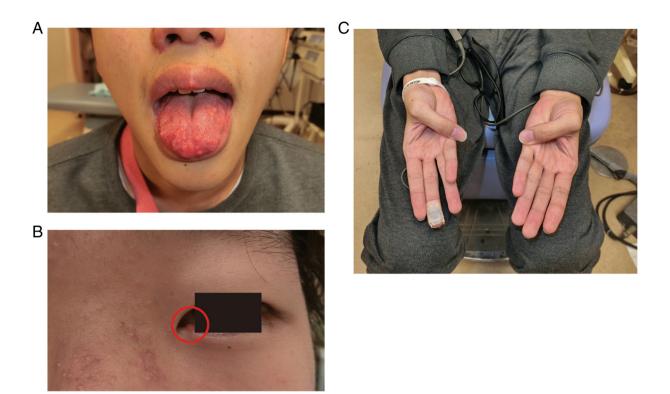


Figure 1. Clinical features of multiple endocrine neoplasia 2B in Case 1. (A) Neuromas of the tongue. (B) A neuroma of an eyelid (surrounded by a circle). (C) Spider fingers (marfanoid habitus).

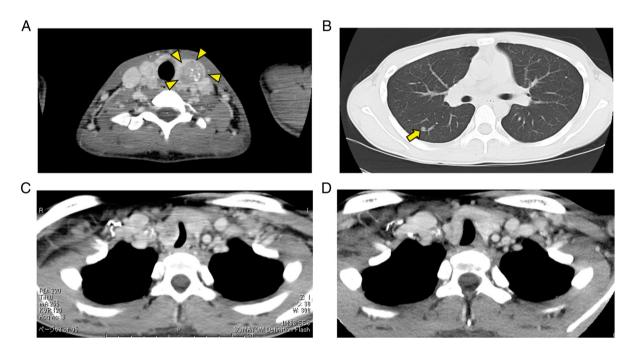


Figure 2. Computed tomography (CT) scan results of Case 1. (A) A thyroid tumor is shown (surrounded by arrowheads). (B) The CT scan demonstrates lung metastasis (pointed with an arrow). (C) The trachea was narrowed and deviated by the thyroid mass. (D) The CT scan shows an improvement in tracheal deviation in Case 1 after initiation of treatment.

was also performed to control electrolyte disturbance-related adverse events due to persistent diarrhea and arrhythmia, such as QT prolongation. Although QT prolongation developed in the patient, the treatment continued without complication. After starting treatment with vandetanib, diarrhea persisted with varying severity. On the contrary, the tumor size was mildly reduced (Fig. 2D). All the nodules in the lungs disappeared. At 25 months after the initial treatment, the patient remains in good condition and the disease has shown no progression. Genetic testing identified the RET mutation in the M918T codon. As the patient confirmed MEN2B, the authors suggested genetic counseling for the family.

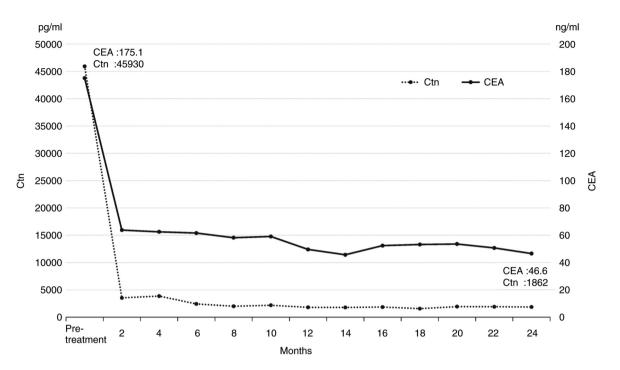


Figure 3. Changes in calcitonin (Ctn) and carcinoembryonic antigen (CEA) levels in Case 1. The levels of Ctn and CEA decreased markedly following the initiation of treatment.

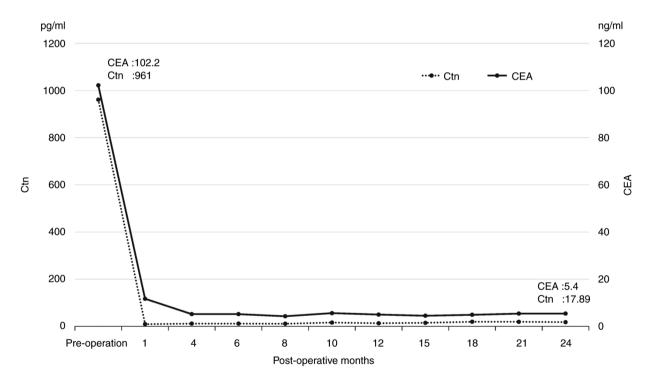


Figure 4. Changes in calcitonin (Ctn) and carcinoembryonic antigen (CEA) levels in Case 3. The levels of Ctn and CEA decreased markedly following surgery.

*Case 2.* The second case was that of the 43-year-old mother of the patient in Case 1. She presented with eyelid neuromas. She was treated for MTC surgically at another hospital when she was 12 years old. Regrettably, the hereditary nature of her disease was not explained to her. It was not until her son was diagnosed as MEB2B that she understood the disease. Genetic testing at Nara Medical University identified her children's RET codon M918T mutation. As her parents had no history of thyroid disease, she may have been the first in her family to acquire an RET codon M918T mutation and developed MTC. Records from her previous hospital showed that she had undergone total thyroidectomy (TT) with bilateral neck dissection and was negative for pheochromocytoma (PHEO). In addition, she had been diagnosed with mucosal neuroma of the tongue based on histopathologic examination. She has been followed up at Nara Medical University and no recurrence or distant metastasis has been observed.

*Case 3.* The third case was that of the 16-year-old elder sister of the patient in Case 1. Although she had no symptoms of MTC, she had tongue neuromas similar to those of her brother. She underwent MTC assessment after her mother and brother were diagnosed. Her blood tests showed abnormally high levels of CEA and Ctn and thyroid masses were detected by ultrasonography. Although an adrenal gland mass was detected by ultrasonography, she was negative for PHEO based on blood test analysis and magnetic resonance imaging. Eventually, MTC was confirmed histocytologically by fine-needle aspiration cytology. Positron emission tomography did not show any signs of lymph node or distant metastases. The patient underwent TT with neck dissection in the central compartment. She exhibits no signs of recurrence 24 months after surgery. Genetic testing also revealed an RET codon M918T mutation.

*Case 4*. The final case was that of the younger sister of the patient in Case 1. She was 10 years old at the time of treatment. She also had neuromas of the tongue and additional ophthal-mologic abnormalities. She had slightly high Ctn levels but no thyroid masses. In addition, no adrenal gland tumors were detected and she was also biochemically negative for PHEO. However, considering the family history of MEN2B, it was decided that prophylactic TT was necessary. Genetic testing identified an RET codon M918T mutation. Postoperative histopathologic examination of the thyroid gland revealed a small amount of MTC tissue. She exhibits no signs of recurrence 21 months after surgery.

# Discussion

MTC is a relatively rare type of thyroid malignancy that arises from parafollicular C cells (1,11). MTC may occur sporadically or may be inherited as a component of MEN 2 syndrome (6,7). RET oncogenes were discovered by Takahashi et al (12) in 1985. All patients with MEN 2 syndrome possess RET mutations. The ATA classifies the risk according to the codon of the gene mutation. Additionally, treatment recommendations are based on the risk category (8). The RET codon M918T mutation, which accounts for 95% of patients with MEN2B, is classified in the highest risk category (4,5,8). TT is recommended for all patients with resectable MTC without evidence of distant metastases. In addition, prophylactic TT for patients with MEN2B associated with RET codon M918T mutation is recommended by the ATA guidelines, even in the first year of life, as they are at high risk for MTC in the future (8).

Although the patients in the present study responded well to treatment, most patients with MTC associated with MEN2B are diagnosed at an advanced state of the disease, as in Cases 1 and 3. In Cases 1 and 3, the levels of Ctn and CEA decreased markedly after their respective treatments (Figs. 3 and 4). The level of Ctn in Case 4 also decreased after surgery although patients with MTC are likely to maintain high Ctn levels after surgery (13).

In Japan, the treatment strategy for asymptomatic MTC with MEN2B is still not standardized. This influenced our

treatment of Case 4, as the patient was asymptomatic but had the gene mutation. We had to obtain approval from the ethics committee and the director of our institution to perform prophylactic TT for Case 4. Due to the lack of available local data, there is still no consensus on the age group on which to perform prophylactic surgery in Japan. In Case 4, postoperative histopathologic examination of the thyroid gland revealed a small amount of MTC tissue and prophylactic TT turned out to be significant.

The patient in Case 2 was not given adequate information about her disease despite undergoing TT for MTC. Therefore, genetic counseling and the lack thereof played an especially important role in our cases. The mother faced difficulty understanding the hereditary nature of their familial disease or the necessity of genetic testing for her children. At first, she was reluctant about genetic testing. However, she gradually understood the significance of testing for RET germline mutations through genetic counseling, which was performed by a multidisciplinary team consisting of head and neck doctors, a medical geneticist, oncologists, nurses and a genetic counselor. Her consent to genetic testing was essential in detecting Cases 3 and 4. Follow-up genetic counseling is required whenever genetic testing is conducted. In 2016, RET genetic testing for patients with MTC received insurance coverage in Japan. However, RET screening before clinical onset is not covered by insurance, even if first-degree relatives have been diagnosed with hereditary MTC. This may result in hesitation to genetic testing and a delay in diagnosis.

In our cases, when the mother was diagnosed with MTC-related MEN2B, there was still insufficient awareness of the genetic characteristics of this disease. Considering this background, the importance of family history and the need for family screening are reaffirmed.

In conclusion, there is no consensus on the age to perform prophylactic TT in Japan. In our case, prophylactic TT prevented MTC from being detected in an advanced state. MEN2B patients should be treated at the optimal time and appropriate genetic counseling is essential in order to prevent delayed diagnosis. We should be aware of the genetic characteristics of MTC-related MEN2B and reaffirm the importance of family history and the need for family screening.

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### Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

# Authors' contributions

AT made substantial contributions to conception and acquisition of data and drafted the manuscript. AT, HU, CM, AN, YY and IO were involved in obtaining the informed consent, literature review, manuscript writing and patient management. TM and TK provided cancer-related scientific inputs and critically revised the manuscript. All authors have accepted responsibility for the entire content of this manuscript and have approved its submission. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

The present study was approved by the Ethics Committee of Nara Medical University Hospital and written informed consent was obtained from the patients.

#### Patient consent for publication

Written informed consent was obtained from the patients for the publication of this case report and any accompanying images.

### **Competing interests**

The authors declare that they have no competing interests.

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