Introduction

On a neurosurgical note, a dumbbell lesion that causes widening of neural foramen and neurological compromise with cord or nerve compression is usually a benign peripheral nerve sheath tumor, such as a schwannoma, or more rarely, an angiolipoma. Ectopic thyroid tissue is an infrequently encountered anomaly caused by abnormal development and disturbed migration of the gland from base of the tongue to the pre-tracheal area in utero (1). Ectopy can often be detected in the tongue, neck, and mediastinal region. The genetic and molecular basis underlying spinal epidural ectopic thyroid carcinoma remains difficult to comprehend, and further discussion of both anatomic and embryological factors is necessary. Only limited reports of ectopic conditions have described malignant transformations of the ectopic tissue. Most of these reports have covered papillary carcinomas. Here, we report a 69-year-old woman with a thoracic epidural mass alongside her spinal cord, which was later confirmed an ectopic follicular variant of papillary thyroid carcinoma (FVPTC).

Case presentation

A 69-year-old woman suffered from persistent left back pain (below the left scapula) for 2 months. An MRI showed a dumbbell mass with contrast enhancement at the T6–7 level and a CT scan showed bone destruction around the lesion (Figures 1, 2). In the local hospital, she underwent a fine-needle aspiration of the lesion and the histopathological report simply showed that collected specimen from spinal lesion was similar to thyroid gland. Thus, she went for our institution for a further diagnosis and treatment. Her neurologic examination demonstrated negative results. She had undergone partial thyroidectomy 15 years prior and histopathology showed thyroid cysts within the right lobe. During follow-up, an ultrasonic examination revealed multiple thyroid nodules graded as TI-RADS 3, and her thyroid function was normal. After hospitalization, she
underwent a decompression and exploration operation, and a soft, extremely vascularized epidural lesion was found. The mass widened the left T6–7 neural foramen and eroded the neighboring osseous tissue. Histopathological results with hematoxylin-eosin (H&E) staining showed ground-glass nuclei with crowding and a grooved appearance (Figure 3). Normal thyroid follicles were found around the neoplastic tissue. In this case, however, papillae were absent, and a follicular architectural pattern was present. Immunohistochemical analysis showed positive TTF-1, Thy, and Ki-67% (index 5%). The patient was diagnosed with an ectopic FVPTC. All of her symptoms were alleviated after surgery.

She developed a worsened soreness in the same site 12 months after the surgical intervention, and her very recent MRI showed further evolvement of the lesion (Figure 4). However, she refused to accept any further treatment.

Discussion

Thyroid ectopia is a developmental disorder that refers to improper location of the gland. Incidence of thyroid ectopia ranges from 0.3 to 1 case per 100,000 population with a female preponderance. The risk of malignant transformation accounts for only 1% of all reported ectopic thyroids (2). Papillary carcinoma is the paramount histologic type in

Figure 1 T2-weighted axial MRI image demonstrating a left epidural mass (white arrow) protruding from the left neural foramen with severe spinal cord compression.

Figure 2 Thoracic CT scan showing enlarged neural foramen and bone destruction.

Figure 3 Histopathological results of the tissue. (A) Photomicrograph showing normal thyroid follicles (original magnification, ×40). (B) Photomicrograph showing a nuclear ground-glass appearance, crowding, and grooving (H&E stain, original magnification ×400).
ectopic tissue, and it is the most commonly encountered thyroid cancer in the normal orthotopic gland. The human thyroid gland originates from thickening of endodermal epithelium, which is termed midline thyroid anlage (MTA), at anterior foregut. Endoderm cells form the anlage and are progenitors of thyroid follicles that later fulfill the central function of the thyroid gland's hormone production. Then cells inside in tandem with adjacent mesenchyme form a diverticulum with a narrow neck that connects the migrating precursor thyroid tissue to the pharyngeal floor (3). The connective duct gradually disappears and the gland expands laterally and descends continuously. Ectopic thyroids develop along the path of migration journey from foramen cecum to the mediastinum and approximately 90% of ectopic thyroid tissue is found in the base of the tongue. However, as the procedure and final completion is under control of interactions between the thyroid itself and neighboring embryonic tissues, a few cases have shown ectopic thyroid glands in the heart, hepatic region, adrenal gland, or ovary (4-7). Some reports covered transformation from parathyroid bodies, whose precursors migrate from the neural crest to thyroid follicles during embryonic life (3).

The terms mass effect and hypothyroidism cover a majority of clinical presentations that lack specificity, and an ectopia faces every problem a thyroid could generate. The relatively benign results of ultrasonic testing (TI-RADS 3) and normal structures of thyroid follicle in the specimen eliminated the possibility of malignant metastasis from the orthotopic thyroid. Thus, our case remains the first concerning ectopic thyroid tissue that developed malignancy and was mistaken for spinal cord neoplasia. The ectopic FVPTC presented as an epidural mass protruding from thoracic neural foramen and finally evolved to a large intra-thoracic mass, which has never been reported before.

Here, we indicate some putative trace of ectopic thyroid originating alongside the spinal cord. (I) Initiative of thyroid genesis is mainly regulated by expression of Titf1/ Nkx2-1, Foxe1, Pax8, and Hhex, and dysregulation of these transcription factors also plays an unequivocal role in thyroid cancer (8). The launch of MTA may be activated by some of the transcription factors, especially Pax8, which is also expressed in correlation with spine and spinal cord formation. (II) In some rare cases, C cell precursors or a portion of them fail to migrate from the neural crest. Instead, they form spinal nerves and other related structures and develop with them (9). Finally, these stubborn cells fail to develop parathyroid bodies but rather form thyroid follicular cells, which has been covered in some reports (3). (III) Spinal ectopia of the gland may be disseminated by errors in the inductive signals required in thyroid migration, and a part of thyroid primordium fails to descend to the settled passage or location.

Thyroid ectopia may happen occultly or with actual or imminent thyroid dysfunction including, in most cases, hypothyroidism and compensatory hyperplasia due to a great occurrence of absent orthotopic thyroid. Diagnosis can be based on physical examination, ultrasonography, or other auxiliary tests. Positron emission tomography/computed tomography (PET/CT) and emission computed tomography (ECT) can help clinicians spot and identify

Figure 4 Post-operative images of the patient. (A) Immediate post-operative MRI showing partial removal of the mass. (B) T2-weighted axial MRI a year after the surgery showing re-growth of the mass.
ectopic glands and evaluate systemic status. CT scan and MRI enable the delineation of soft tissues with contrast. Optimal treatment modalities for an ectopic thyroid depend on tumor site and size, patient age and sex, comorbidity, hormone status, and other correlating factors. An asymptomatic ectopia or euthyroid condition warrants no treatments (10). Usually, levothyroxine replacement suffices for cases of hypothyroidism alone. However, when a patient complains about the mass effect in their body parts combined with severe thyroid dysfunction or the ectopic gland show signs of malignant transformation, radical interventions are necessary. As with normal thyroid tumors, a surgical approach and radioiodine therapy are often the best way to remove the lesion and are considered the standard of care (11,12). It is necessary to examine the orthotopic thyroid via cervical ultrasonography before total surgical resection to prevent permanent hypothyroidism.

We here report a case of ectopic thyroid mimicking a spinal dumbbell tumor, which has never been described before. A benign peripheral nerve sheath tumor or a metastasis from thyroid should be considered in patients in whom there is no reason to suspect spinal ectopia. Mechanisms underpinning ectopic thyroid remain to be fully determined and those underlying ectopia growing both within and outside the spinal canal are likely to be multifactorial.

Acknowledgments

We thank LetPub (www.letpub.com) for its linguistic assistance during the preparation of this manuscript.

Footnote

Conflicts of Interest: The authors declare no potential conflicts of interest.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. Written informed consent was obtained from the patient for publication of this Case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

References

2. Jensen PH, Pilt AP, Wittendorff HE. Ectopic thyroid tissue with a primary carcinoma. Ugeskr Laeger 2017;179: