Role of immunohistochemistry in metastatic clear cell variant of follicular thyroid carcinoma: A case report

Yash Pradeep Vaidya, Rajan Vaithianathan, Ramanathan Manickam, Dhananjay Kotasthane

ABSTRACT

Introduction: Clear cell variant of follicular thyroid carcinoma with synchronous bony metastasis and a normal thyroid stimulating hormone level, is an extremely rare condition.

Case Report: A 55-year-old male was presented to us with a painful swelling in the right arm. The biopsy showed clear cell adenocarcinoma, raising concerns for a metastatic renal cell carcinoma. Computed tomography scan of abdomen failed to show any renal lesions. A detailed physical examination revealed a small nodule of the right thyroid lobe. Fine needle aspiration cytology (FNAC) of the nodule was reported as follicular neoplasm. A right hemithyroidectomy and the subsequent completion thyroidectomy showed clear cell type of follicular thyroid carcinoma. Immunohistochemistry for thyroglobulin (Tg) further confirmed the diagnosis.

Conclusion: Clear cell variant of follicular thyroid carcinoma is a very rare condition, hence a high index of suspicion is essential for diagnosis. The importance of performing a detailed physical examination cannot be more emphasized as small thyroid lesion like in this case can be easily missed, leading to a delay in diagnosis.
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Keywords: Clear cell variant, Follicular thyroid carcinoma, Thyroglobulin, Thyroidectomy

INTRODUCTION

Follicular thyroid carcinoma is the second most common type of thyroid cancer, comprising around 10–15% of thyroid malignancies. It has been histologically classified into oncocytic and clear cell types by the World Health Organization [1]. Follicular thyroid carcinoma with clear cell change is rare, and only some cases have been reported in literature [2]. This can be a diagnostic problem as metastatic clear cell carcinoma to the bones is usually attributed to a renal primary. Moreover, the most common cause for metastatic lesion in the thyroid is also renal cell carcinoma [3]. This clinical scenario may prove very challenging in arriving at an accurate diagnosis and for optimum management.
CASE REPORT

A 55-year-old male initially presented with pain and difficulty in moving his right arm after a fall. No significant past medical or family history was noted. On examination, he had a hard, 6x4 cm swelling over the upper third of the humerus. X-ray of the right humerus showed a soft tissue mass and a lytic lesion causing pathological fracture with varus angulation. Magnetic resonance imaging scan showed an intramedullary lesion in the bone with soft tissue extension, suggestive of a primary bone malignancy (Figure 1).

An open biopsy of the lesion revealed clear cell adenocarcinoma, probably of renal origin (Figure 2). However, immunohistochemistry staining using vimentin, CD 10, cytokeratin and S100 were equivocal. Computed tomography scan of abdomen was normal (Figure 3). On further examination, a small nodule was palpable in the right lobe of the thyroid. Thyroid function tests were within normal limits. Ultrasound of the neck demonstrated a 2x1 cm, hypoechoic nodule in the right lobe. Ultrasound guided FNAC was suggestive of a follicular neoplasm. A U-slab was applied for the fracture in right humerus.

The patient underwent an initial right hemithyroidectomy. Preoperatively, the patient was explained in detail about the need for this procedure as well as for completion thyroidectomy if malignancy was confirmed. Histopathology proved it to be clear cell variant of follicular thyroid carcinoma, with tumor cells invading into vascular spaces, features very similar to the biopsy from the right humerus (Figure 4). He subsequently underwent completion thyroidectomy to remove the left lobe and this also showed similar features.

Immunohistochemistry was strongly positive for thyroglobulin and negative for prior mentioned markers, thus confirming the lesion to be a thyroid primary rather than a metastatic lesion from the kidneys.

The patient made an uneventful recovery, with no symptoms of hypocalcaemia. He was later referred to the regional Radiation Oncology Centre for radioactive iodine (RAI) treatment and planned for a RAI scan (both thyroid and whole body). We plan to monitor the patient regularly with serum thyroglobulin levels.

DISCUSSION

Most follicular thyroid carcinomas are asymptomatic and diagnosed late. Thyroid scintigraphy usually shows a cold nodule. They typically present as solitary lesions, and very rarely involve both the lobes. Distant metastasis is common when there is capsular or vascular invasion [1]. Follicular thyroid carcinoma commonly spreads by the hematogenous route. Around 10–15% of patients have distant metastasis, occurring more often in tumors larger than 2 cm size [4]. The most common sites of distant metastases are the bones (with typical lytic lesions).
and lungs followed by the brain, liver, bladder and skin [5]. The incidence of metastatic disease is higher in the oncocytic variant than the clear cell type [6].

The clear cell appearance has been attributed to either accumulation of vesicles or glycogenation, thyroglobulin accumulation and hypertrophy of the Golgi complex due to excessive thyroid stimulating hormone stimulation [1, 7, 8]. Presence of clear cell features in follicular thyroid carcinoma very rare in patients with a normal thyroid stimulating hormone value as noted in this patient.

Almost 25–30% patients with renal cell carcinoma have distant metastasis at the time of diagnosis [9]. Hence this is usually the first differential diagnosis in such cases.

When metastatic lesions in the bone and the thyroid exhibit clear cell changes, it becomes difficult to differentiate metastatic renal cell carcinoma from a primary clear cell follicular thyroid carcinoma [10]. In such cases, immunohistochemistry plays an important role in clinching the origin of the tumor cells, as we found in our case. Clear cell renal cell carcinoma usually stains positive for vimentin, cytokeratin antibodies (AE1/AE3), CD10, renal cell carcinoma marker (RCCM), PAX2, PAX8, and carbonic anhydrase IX (CAIX), and stains negative for cytokeratins (CK7, CK 20), high-molecular-weight cytokeratin (HMWCK), CD 117 and parvalbumin [11].

CONCLUSION

Clear cell variant of follicular thyroid carcinoma is a very rare condition, hence a high index of suspicion is essential for diagnosis. Metastatic carcinomas exhibiting clear cell changes are commonly mistaken for a spread from a renal cell carcinoma. The importance of performing a detailed physical examination cannot be more emphasized as small thyroid lesion like in this case can be easily missed, leading to a delay in diagnosis. Immunohistochemistry is the definitive way of differentiating clear cell carcinomas arising from thyroid and the kidneys.

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Author Contributions

Yash Pradeep Vaidya – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Rajan Vaithianathan – Substantial contributions to conception and design, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Ramanathan Manickam – Substantial contributions to conception and design, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Dhananjay Kotasthane – Substantial contributions to conception and design, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor

The corresponding author is the guarantor of submission.

Conflict of Interest

Authors declare no conflict of interest.

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REFERENCES


ABOUT THE AUTHORS


**Yash Vaidya** is Final Year Resident at Department of General Surgery, Mahatma Gandhi Medical College and Research Institute, Pondicherry, India. He has published multiple research papers in national and international academic journals. He intends to pursue career in cardiothoracic and vascular surgery in future.

**Rajan Vaithianathan** is Associate Professor at Department of General Surgery, Mahatma Gandhi Medical College and Research Institute, Pondicherry, India. He has published several research papers in national and international academic journals. His field of interest is surgical gastroenterology.

**Ramanathan Manickam** is Professor and Head at Department of General Surgery, Mahatma Gandhi Medical College and Research Institute, Pondicherry, India. He has published several research papers in national and international academic journals. His field of interest is surgical gastroenterology.

**Dhananjay Kotasthane** is Professor and Head at Department of Pathology, Mahatma Gandhi Medical College and Research Institute, Pondicherry, India.
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