Angiosarcoma of the Thyroid Gland: A Case Report

Fouad Zouaidia\textsuperscript{a, b}, Ahmed Jahida, Hafsa Elouazzanica, Leila Larakia, Zakiya Bernoussica, Fatima Mansouri\textsuperscript{a}, Najat Mahassini\textsuperscript{a}

Abstract

The authors described a case of rare histological type of primary thyroid tumour: angiosarcoma. We report a case of 37 years old female who presented with the mass in the anterior of the neck that had been rapidly growing for several months. Scintigraphy showed a cold nodule in the left lobe of the thyroid. The patient underwent subtotal thyroidectomy. The cut surface showed a bulging solid grey hemorrhagic mass, measuring 6.5 cm at the lower pole of the left lobe. Microscopically a poorly differentiated highly desmoplastic polymorphocellular tumour. Immunohistochemically, the cells showed immunoreactivity for endothelial markers CD31, CD34, Factor VIII related antigen and immunonegativity for epithelial markers including pancytokeratin, epithelial membrane antigen and thyroglobulin. Radiotherapy was applied. The patient was alive after a follow up of five months.

Keywords: Thyroid; Angiosarcoma; Pathology

Introduction

Angiosarcoma, made up of malignant endothelial structures, represents a rare and aggressive connective tissue tumour of the thyroid. It rarely occurs in that organ and most of these occur mainly in patients from the mountainous Alpine regions. It is most prevalent in Switzerland where it constitutes 4\% to 5\% [1].

Angiosarcoma of the thyroid has long been a controversial entity because some authors believe that it is a distinct entity of endothelial origin but others consider it to be a variant of undifferentiated carcinoma. So the differential diagnosis includes pseudoangiosarcomatous carcinoma of the thyroid [2].

We describe a case of angiosarcoma of the thyroid. To confirm the nature of the present tumour, immunohistochemistry examination was done.

Case Report

A 37 years old female presented with the mass in the anterior of the neck that had been rapidly growing for several months. Scintigraphy showed a cold nodule in the left lobe of the thyroid. The results of thyroid function tests were within the normal range.

The patient underwent subtotal resection of the left thyroid lobe. The cut surface showed a bulging solid grey hemorrhagic mass, measuring 6.5 cm at the lower pole of the left lobe (Fig. 1). The specimen was fixed in 10\% neutral formalin and then embedded in paraffin. Microscopically it was a poorly differentiated highly desmoplastic polymorphocellular tumour (Fig. 2, 3).

Immunohistochemistry was done by avidin biotin peroxidase complex method. Antibodies against the following antigens were used. Immunohistochemically, the cells showed immunoreactivity for endothelial markers CD31, CD34 (Fig. 4),...
Facor VIII related antigen and immunonegativity for epithelial markers including pancytokeratin, epithelial membrane antigen and thyroglobulin.

Radiotherapy was applied. The patient was alive after a follow up of five months.

Discussion

Angiosarcoma is uncommon soft tissue neoplasm that accounts for less than 1% of all sarcoma. Angiosarcoma involving thyroid is a rare entity, more often described in the Alpine region [3]. The location in the thyroid gland is nevertheless quite exceptional.

The sex distribution was female predominant (female-male ratio 9:3), the age at diagnosis ranging from 50 to 88 years. The typical history is a sudden and rapid increase in the size of long standing goitre. Common signs and symptoms besides the neck swelling include hoarseness, radiation pain, dysphagia, and marked loss of weight. The presence of goitre was stated in our patient. Progression into the mediastinum with compressive complications has also been described [4].

The tumour ranges in size from 2.5 to 11 cm and generally occurs singly. The cut surface of the thyroid shows a well demarcated bulging, variated and hemorrhagic dark red lesion.

On histologic examination, this tumour typically discloses anastomosing capillary-sized or dilated spaces lined by large, atypical endothelial cells and containing red blood cells. The endothelial differentiation is shown by immunoreactivity for endothelial markers such as CD31, CD34, Ulex europaeus, factor VIII related-antigen and immunonegativity for epithelial markers including pancytokeratin, epithelial membrane antigen and thyroglobulin. However, a rare aberrant expression of cytokeratin in angiosarcoma in other organs has occasionally been reported because embryonic endothelial cells may express keratin [5, 6].

Very often, the tumour shows a well developed fibrous capsule, as seen in our case with remnants of thyroid follicles scattered among the tumour tissue and compressed against the capsule. These features strongly suggest that the tumour has developed on top of goitrous nodule or an adenoma.

The diagnosis is mostly difficult and the histological features mimicking anaplastic carcinoma of the thyroid and other sarcoma.

Focal solid undifferentiated areas are invariably present consisting of spindle cells arranged in fascicular pattern reminiscent of the most common morphologic pattern encountered in an anaplastic carcinoma [7, 8]. This latter feature make many experienced pathologists unfamiliar with this lesion to regard it as a variant of anaplastic carcinoma or other sarcoma. In such circumstances, angiosarcoma can be distinguished from it by immunohistochemistry. To date, CD31 is considered the most sensitive and specific marker for endothelial differentiation, being expressed in 90% of angiosarcoma and in less than 1% of carcinomas [3, 9].

Its etiology remains unknown. The prognosis of this tumour is worst because it is growing very fast and can develop metastases. These are found most frequently in the lungs and cervical or tracheobronchial lymph nodes [1].

Therapy is a challenge with frequent difficulties. Surgery represents the first step of choice in cases of radically removable tumour as seen in our patient. Radiation therapy may be effective in some patients and can be completed us-
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Conflict of Interest

No competing interest.

References