

Low Back Pain as the First Presentation of Hürthle Cell Carcinoma of Thyroid

Lombalgia como a primeira manifestação do carcinoma de células de Hürthle

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Abstract

Hürthle cell carcinoma is a rare variant of differentiated thyroid cancer and occasionally generates distant metastases. There are few reports of Hürthle cell carcinoma metastases to the vertebral column and none with sacral involvement. Here, we report a 48-year-old woman with a 2-year history of moderate low back pain, alleviated with analgesic medication. Her previous medical files were unremarkable. She performed medical evaluation, and a magnetic resonance of lumbar spine revealed a solitary hypointense T2 lesion on the left lamina of S1. The patient was submitted to percutaneous biopsy that displayed carcinomatous pattern compatible with Hürthle cell carcinoma of thyroid. After that, she was submitted to thyroid ultrasound that showed a nodule. Even in the presence of metastases, patients with Hürthle cell carcinoma usually have a relatively good prognosis. We discuss aspects of the diagnosis, management, and surgical treatment of metastatic Hürthle cell carcinoma in reference to the literature.

Keywords

- ▶ low back pain
- ▶ spinal diseases
- ▶ neoplasm metastasis

Resumo

O carcinoma das células de Hürthle é uma variante rara do câncer de tireoide. Existem poucos relatos de metástases dessas lesões para a coluna vertebral. Nós relatamos o caso de uma paciente de 48 anos com história de dois anos de lombalgia. Ela era previamente hígida. A avaliação radiológica evidenciou uma lesão solitária na lamina esquerda de S1. Após biópsia percutânea, foi diagnosticado o carcinoma, sendo diagnosticada também uma lesão sincrônica em tireoide. Nós discutimos os aspectos, clínicos, radiológicos e de tratamento no caso supracitado.

Palavras-chave

- ▶ dor lombar
- ▶ doenças da coluna vertebral
- ▶ metástase neoplásica

Introduction

Thyroid cancer is the most common tumor of the endocrine system.^{1,2} It is usually divided in three major subtypes: papillary, follicular, and medullary. The prognosis of patients

with differentiated thyroid cancer is excellent and 10-year survival is more than 90%.¹ Dissemination of thyroid cancer is primarily via regional lymph nodes and metastatic pulmonary disease may develop, followed by bone involvement.¹

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Hürthle cell carcinoma is a rare variant of differentiated thyroid cancer and occasionally generates distant metastases.²⁻⁴ There are few reports of Hürthle cell carcinoma metastases to the vertebral column and none with sacral involvement.²⁻⁴ We report a unique case of Hürthle cell carcinoma metastasis to sacral spine presenting with low back pain.

Case Description

We report a 48-year-old woman with a 2-year history of moderate low back pain irradiating to posterior and medial left thigh, being initially alleviated with common analgesic medication and physical therapy. After 1 year, she returned in use of paracetamol and codeine with a milder sustained pain that increased at walking or long standing up.

Her previous medical files were unremarkable. At that moment, she denied fever or weight loss. She underwent new medical evaluation. Her physical examination, including neurologic status, was normal. Because of long course of pain, she was submitted to radiologic evaluation. Lumbar and sacral spine radiographs were initially of limited help,

but a lumbar and sacral tomography immediately revealed an osteolytic lesion in the left S1 lamina, invading the vertebral canal. A magnetic resonance imaging revealed a solitary hypointense T1 and T2 lesion on the left lamina of S1, which homogeneously enhanced after endovenous gadolinium injection (►Fig. 1). Among possible diagnosis, tumoral etiology was considered, and because of topography and image, the main hypothesis included sacral chordoma, lymphoproliferative neoplasms, and metastatic lesions. To determine the nature of lesion, the patient was submitted to percutaneous biopsy guided by fluoroscopy, without any surgical complications.

Pathological findings were compatible with an oncocytic solid-glandular neoplasm. Immunohistochemistry showed expression of tireoglobulin, CK7, and CK19, a pattern compatible with Hürthle cell carcinoma of thyroid (►Fig. 2). Immediately after diagnosis, she was submitted to thyroid ultrasound that revealed a solitary, solid, lobulated, heterogeneous, predominantly hypoechoic with hyperechoic foci nodule, with peripheral and central vascularization in the middle third of right thyroid lobule. She was forwarded to Neck and Head Surgery Service for further approach.

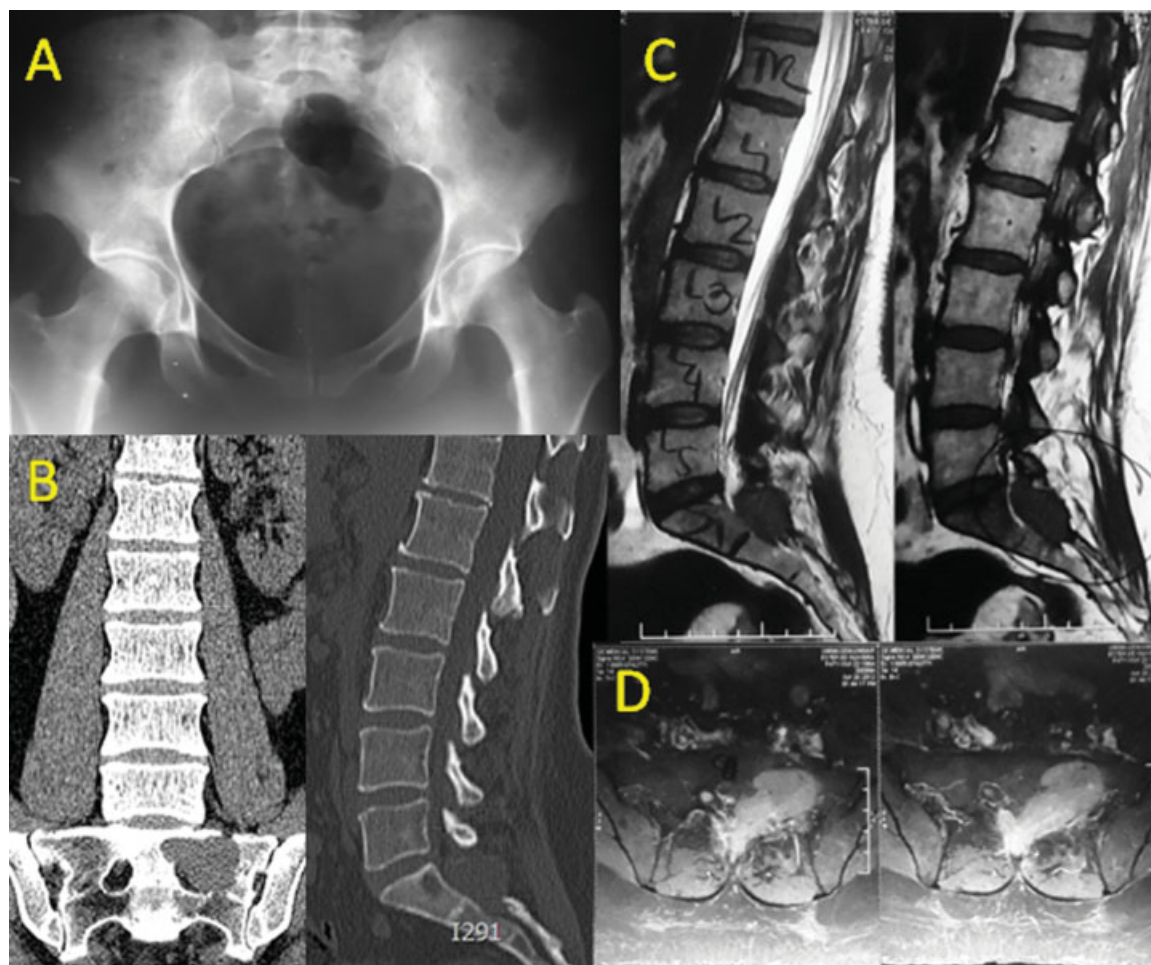


Fig. 1 Radiologic evaluation. (A) A pelvic radiography with interference of gas in the colon. (B) Lumbar and sacral tomography revealing an osteolytic lesion in left S1 lamina, invading vertebral canal. (C) Sagittal magnetic resonance imaging displaying a solitary hypointense T1 and T2 lesion on the left lamina of S1. (D) Axial magnetic resonance imaging on the left lamina of S1 homogeneously enhancing after endovenous gadolinium injection.

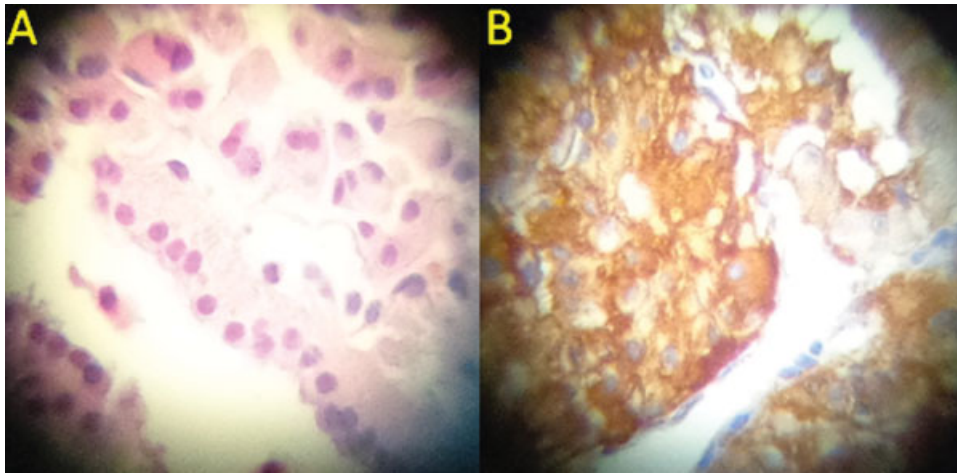


Fig. 2 Pathological findings. (A) Histologic features of thyroid carcinoma in hematoxylin-eosin staining. (B) Immunohistochemistry revealing intense tireoglobulin staining, a typical finding of Hürthle cell carcinoma.

Discussion

Thyroid cancer bone metastases frequently cause significant morbidity.^{1,2} They may be present at the initial diagnosis in up to 50% of patients.² Spread of thyroid carcinoma to bone is more common in patients older than 45 years and is characteristically symptomatic and multicentric.² In one series, the vertebrae, pelvis, ribs, and femur were respectively the most common sites of metastases. Multiple lesions were present in 53% of the cases. The overall 10-year survival rate from the time of diagnosis of thyroid cancer was 35%, and from diagnosis of initial bone metastasis was 13%.²

Hürthle cell carcinoma is a rare variant of differentiated thyroid cancer characterized by solid tumor nests and microfollicle formation, sometimes with abundant colloid production and strong immunocytochemical reactivity for thyroglobulin.^{3,4} Even in the presence of metastases, patients with Hürthle cell carcinoma usually have a relatively good prognosis, being considered the most favorable histologic subtype for survival.³

Few other reports disclosed Hürthle cell carcinoma as spinal metastasis, and none with sacral involvement.³⁻⁵ We highlight our case once it approaches a young woman

presenting with atypical symptomatology of low back pain as first clinical presentation of a solitary sacral bone metastatic Hürthle cell carcinoma.

Conflict of Interest

The authors declare no conflicts of interest.

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