Erdheim-Chester Disease: Case Report with Aggressive Multiple Organ Manifestations

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Abstract

Erdheim-Chester disease (ECD) is a rare non-Langerhans cell, lipid-laden histiocytosis with specific histological and radiological findings. The diagnosis sometimes is established lately in the course of the disease. We present a case of a 64-year-old female with elevated inflammatory markers for one year and symptoms related with her comorbidities, particularly bone pain and short of breath. Past medical history includes a stage III chronic kidney disease, central diabetes diagnosed when she was 38 years old, Paget Disease, metabolic syndrome and ischemic cardiopathy. Computed tomography in the past showed a tissue densification in the thoracic vertebal column and kidneys with hairy aspect. X-ray of the arms, legs, skullcap, and demonstrated sclerotic changes. F-Fluorodeoxyglucose positron emission tomography showed uptake in the skull, mediastinum, abdomen and long bones from arms and legs. Biopsy of the hairy kidney was consent after 4 years of an unknown disease in progression. Histological findings of the biopsy reported a diffuse infiltration by foamy histiocytes. On immunohistochemical staining, the histiocytes were positive for CD68 and negative for CD1 and S100. Mutation of BRAF V600E was present and ECD was established. Tocilizumab was initiated off label due to psychiatric contra indication for interferon use and no clinical conditions for BRAF inhibitors and symptoms started being controlled.

The diagnosis of ECD is usually challenging due to the rarity of the disease and clinical overlapping with many other conditions. The rarity and variable presentation of this disease usually leads to delayed diagnosis and to high morbidity and mortality rates from associated complications.

Key Words: non-langerhans cell; multiple organ; braf v600e

Learning Points

Rarity of this disease makes the diagnosis challenging and to a high morbidity and mortality rates from associated complications.

1. New treatments are being available and this case report shows a good response to a novel therapeutic that is not fully approved to the disease.

Case Description

Introduction

Erdheim-Chester disease (ECD) is a rare non-Langerhans cell, lipid-laden histiocytosis with specific histological and radiological findings. [1] Its estimated there are 600 cases described in the literature. The disease usually presents between 40 and 60 years and its etiology is still unknown but BRAF proto-oncogene has been identified in the majority of ECD cases.

The diagnosis of ECD is made with a combination of clinical presentations and imaging features, that may be present on other diseases, that is why the diagnose sometimes is established lately in the course of the disease. Heterogenous clinical manifestations depend on site and degree of involvement, but around 95% of ECD patients have skeletal involvement and bone pain is the main symptom. Apart from skeletal involvement, ECD may infiltrate eyes, endocrine organs, kidney, heart and central nervous system (CNS). [2]

The differential diagnosis of ECD includes diseases such as Langerhans cell histiocytosis, multiple sclerosis, Paget disease, amyloidosis and others.

The first line treatment is interferon alfa although it might not be eligible for all the patients depending of the comorbidities and efficacy is limited against severe manifestations, specifically CNS and cardiovascular involvement. Treatment with other agents is based on case reports described in the literature.

In this paper, we present an ECD case with multiple organs involved because of the delay between the initial symptoms and the diagnosis. To the best of our knowledge, this is one of the few cases reported from Portugal.

Case report

A 64-year-old female presented with elevated inflammatory markers for one year and symptoms related with her comorbidities, particularly bone pain and short of breath. Past medical history includes a stage III chronic kidney disease, diabetes insipidus diagnosed when she was 38 years old, Paget Disease, metabolic syndrome and ischemic cardiopathy. On admission to our appointment, the patient’s general appearance was
normal and hemodynamically stable. The initial laboratory workup revealed mild anemia (11.2 g/dL), normal white blood cell count and normal platelets. Chronic kidney disease was supported with creatinine of 1.6 mg/dL. Liver function was normal. The inflammatory markers such as C-reactive protein (CRP) and the erythrocyte sedimentation rate (ESR) were elevated - CRP: 106 mg/dL (reference values <5.0 mg/dL); ESR 93 mm (reference values <20 mm). Patient had made a computed tomography (CT) in the near past that showed an anterior tissue densification in the thoracic vertebral column and kidneys with hairy aspect (Figure 1).

The patient underwent radiological studies, with bilateral (Figures 2 and 3).

F-fluorodeoxyglucose positron emission tomography (FDG-PET) showed uptake in the skull, mediastinum, abdomen and long bones from arms and legs (Figure 4).
Cardiac MRI revealed a fibrotic structure next to the free wall of the right atrium. After these results we started questioning the past medical history of Paget Disease and diabetes insipidus and thought these findings probably would come from the same disease. A kidney and a bone biopsy were proposed but patient denied it for 4 years until she was admitted to the ward because of a cardiac dysthymia (needing definitive pacemaker) with decompensated heart failure, aggravated pericardium effusion and acute on chronic kidney failure. Surgery was proposed for intraperitonealization of the ureters and biopsy of the hairy kidney was consent after 4 years of an unknown disease in progression. Histological findings of the biopsy reported a diffuse infiltration by foamy histiocytes. On immunohistochemical (ICH) staining, the histiocytes were positive for CD68 and negative for CD1 and S100. Mutation of BRAF V600E was present and ECD was established. Once the diagnose of ECD was confirmed and because of its aggressive nature and involvement in this patient treatment was started immediately. She had a psychiatric contra indication for interferon use and had no clinical conditions for BRAF inhibitors. She started treatment with tocilizumab 8mg/kg/month intravenous based on its use described in case reports. Our patient had a good response after starting tocilizumab. Until now she is under surveillance with a stable disease.

**Conclusion**

ECD is a rare, multi system histiocytic neoplasm characterized by tissue infiltration or organ encasement by CD68+, CD1+, S100-low/negative foamy histiocytes. Heterogenous clinical manifestations depend on site and degree of involvement. Skeletal involvement with sclerotic lesions and radiotracer uptake is common. Other disease manifestations include neurologic symptoms due to CNS involvement, diabetes insipidus, constitutional symptoms and heart and lung involvement. When affected, CNS, lungs, heart and retroperitoneal space account for a severe prognosis. [3]

In our patient diabetes insipidus was diagnosed when she was 38 and it might be the first manifestation of ECD, misinterpreted for almost 30 years. The diagnosis of ECD is usually challenging due to the rarity of the disease and clinical overlapping with many other conditions. In our case patient was being followed in orthopedic appointment because of bone lesions thought to be related with a Paget Disease. The rarity and variable presentation of this disease usually leads to delayed diagnosis and to high morbidity and mortality rates from associated complications. With the progression of the disease until the diagnose is established sometimes patients may not be eligible for first line treatments, as occurred with our patient where we used tocilizumab based on case reports.

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**References**


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