We report a case of mediastinal granulocytic sarcoma with cardiac infiltration in a young man with no evidence of leukemia involving the bone marrow or peripheral blood. Diagnosis was accomplished by immuno-histochemistry with expressions of myeloperoxidase and CD99 antigens. The patient achieved clinical remission, but evolved with febrile neutropenia during chemotherapy and died. Although subclinical cardiac infiltrations are commonly found at autopsy in patients with acute non-lymphoblastic leukemia, only one case of involvement of the heart with granulocytic sarcoma in the absence of bone marrow disease has been published in the literature. A diagnosis of granulocytic sarcoma should not be excluded when the biopsy of the bone marrow does not show any evidence of leukemic infiltration. Rev. bras. hematol. hemoter. 2008; 30(4):339-341.

Key words: Granulocytic sarcoma; leukemia; cardiac infiltration.

Introduction

Granulocytic sarcoma (GS) is a rare extramedullary tumor composed of immature myeloid cells. It is usually associated with leukemia or other myeloproliferative disorders but can also occur without overt hematologic diseases. GS can occur before, concomitantly, or after the overt development of acute or chronic myelogenous leukemia. Although this tumor is known to occur in almost every site of the body, cardiac involvement and a mediastinal mass are rare. The involvement of the heart with GS with no evidence of leukemia involving bone marrow or peripheral blood is very rare and usually diagnosed at autopsy. Only one report has been published in the literature. Histologic testing is essential to provide the proper diagnosis.

We report the clinical and pathological features of a mediastinal GS with cardiac infiltration in a young adult man with no evidence of leukemia involving bone marrow or peripheral blood.

Case Report

A 31-year-old man with a five months history of dry cough and progressive dyspnea was admitted to our hospital due to pain in the left hemithorax when he was in dorsal decubitus. Physical examination showed vesicular murmur decreased in the right lung and irregular heart rhythm with a third heart sound. Patient had done previously echocardiography, which revealed mass of homogeneous texture of 58mm x 54mm, occupying approximately 65% of the left atrium, which is highly increased. The mass invades the cavity of the left ventricle (VE) in diastole through the mitral valve, which caused great restriction on the filling of the VE. He was submitted to surgery with diagnostic hypothesis of atrial myxoma. However, it was not held resection because the lesion was too extended. The diagnoses prior morphological and immunohistochemistry were angiosarcoma of low malignancy, paraganglioma heart and small cell carcinoma. Directed chemotherapy treatment was started with Etoposide and Cisplatin, with no good clinical
Immuno-histochemistry: A - Myeloperoxidase; B - MIC - 2 expression by neoplastic cells (ABC x100).

Discussion

Granulocytic sarcoma is usually associated with leukemia or other myeloproliferative disorders, but can also occur without overt hematologic disease, i.e. in patients with a normal bone marrow and no history of acute myelogenous leukemia. Although cardiac infiltration is common in advanced stage of acute leukaemia, it is not usually diagnosed at life. But in this case the cardiac involvement was detected when the patient was still alive. The transthoracic echo revealed mass of homogeneous texture occupying the left atrium. Our research in literature disclosed the presence of only one case of mediastinal nonleukemic SG with cardiac infiltration.

GS poses a diagnostic pitfall for both pathologists and oncologists. Previous literature showed that almost half of the patients with primary GS were misdiagnosed initially. Careful attention should be paid to the relatively high incidence of hematologic malignancies in mediastinal tumors. Early and accurate diagnosis of these tumors is essential because some of these patients require immediate treatment by hematology specialists.

This case was originally misdiagnosed as angiosarcoma of low malignancy, paraganglioma heart and small cell carcinoma, which resulted in inappropriate therapy. The positivity for myeloperoxidase and MIC-2 (CD 99) at immunohistochemistry indicated the correct diagnostic of GS. Directed chemotherapy treatment was started with Daunorubicina and Cytarabine, with clinical remission. Histopathologists should be aware that GS might occur in unusual extramedullary sites without evidence of bone marrow involvement.

Resumo


Palavras-chave: Sarcoma granulocítico; leucemia; infiltração cardíaca.

References


Avaliação: Editor e dois revisores externos
Conflito de interesse: não declarado
Recebido: 14/02/2008
Aceito 16/04/2008