Clinical case

Diagnosis of pancreatic plasmacytoma by endoscopic ultrasound-fine needle aspiration

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Abstract

Plasmacytoma presents more frequently in middle age men with aerodigestive tract involvement, especially in the head and the neck. Gastrointestinal tract involvement is uncommon, but the organ most commonly involved is the stomach. We report the first case in the literature in which final diagnosis was made by fine-needle aspiration biopsy guided by endoscopic ultrasound with adequate sample for pathologic analysis. The treatment of this entity is systemic chemotherapy but its effectiveness is limited. Plasmacytoma should be taken into account in differential diagnosis of pancreatic masses.

Key words: Plasmacytoma, neoplasms, endoscopic ultrasound, fine-needle aspiration, pancreatic masses, Brazil.

Resumen

El grupo de pacientes con la mayor incidencia de plasmocitomas está compuesto por hombres de edad media, con involucro del tracto aerodigestivo, especialmente en cabeza y cuello. La afectación del tracto gastrointestinal no es común; el órgano afectado con mayor frecuencia es el estómago. Presentamos en primer caso en la bibliografía en el que el diagnóstico definitivo fue hecho mediante una punción con aguja fina guiada por ultrasonido endoscópico, lo que permitió obtener un adecuado material para el análisis histopatológico. La única posibilidad de tratamiento es la quimioterapia sistémica, pero su efectividad es limitada. El plasmocitoma debe ser considerado en el diagnóstico diferencial de masas pancreáticas de origen desconocido.

Palabras clave: Plasmocitoma, neoplasias, ultrasonido endoscópico, aspiración con aguja fin, masa pancreática, Brasil.
Introduction

The solitary plasmacytoma usually affects the bone tissue, but its extramedullary form, with incidence around 5%, may be related to secondary invasion by multiple myeloma. The primary form is extremely rare. We report the first case in the literature of pancreatic plasmacytoma diagnosed by ultrasound-guided fine needle aspiration (EUS-FNA).

Case report

Forty-two years-old male patient, with diagnosis of solitary plasmacytoma in the right shoulder since 2002, was treated initially with local radiotherapy with resolution of the lesion. In December 2004, new plasmacytomas in knee and ankle were diagnosed and were treated with radiotherapy and steroids (dexamethasone). Good response was achieved, but in November 2007 a new plasmacytoma in the right thigh was diagnosed, and lambda light chain was detected in the urine. At this time, there were no target organ injuries or bone marrow plasmacytosis, but because of the multiple plasmacytomas systemic treatment was chosen. This treatment was conducted with radiotherapy and chemotherapy (cyclophosphamide, dexamethasone, thalidomide - seven cycles), without response. A new chemotherapy protocol with bortezomib, dexamethasone and thalidomide was started. After completing the second chemotherapy cycle of this protocol, the patient developed cholestatic jaundice and was admitted for investigation. A CT scan of the abdomen showed a large expansive lesion in the head of the pancreas (Figure 1). Patient underwent ERCP to clear the biliary tree and EUS was performed. This study showed hypoechoic and heterogeneous mass in the head of the pancreas. FNA was performed and samples were obtained for pathologic analysis (Figure 2). The cytology report revealed small cell neoplasm with immunohistochemical profile consistent with plasmacytoma producing lambda light chain (Figure 3).

Chemotherapy protocol was changed to bortezomib, dexamethasone and cyclophosphamide. Four cycles of chemotherapy were administered with partial response. Patient died because infectious complications after the fourth cycle of chemotherapy.

Discussion

Extramedullary plasmacytomas are malignant tumors of monoclonal plasma cells. Progression to multiple myeloma is seen in approximately 11 to 30% of cases within 10 years. The group with highest incidence is composed of middle age men with aerodigestive tract involvement, especially in the head and the neck. The gastrointestinal tract involvement is uncommon, but the organ most commonly involved is the stomach. Rarely, the pancreas is infiltrated by plasmacytes, either as solitary plasmacytoma or extramedullary invasion by multiple myeloma. The diagnosis is based on the finding of extramedullary monoclonal plasma cells without proliferation of plasmacytes in bone marrow.

Computed tomography (CT) is the most widely used method for assessing these cases. The most frequent location is the head, but there are reports of diffuse involvement and two plasmacytomas in the same pancreas.

The CT-guided fine needle aspiration can make the diagnosis. Due to the similar cytomorphology of the plasmacytes in relation to neuroendocrine cells, pancreatic extramedullary plasmacytoma

Figure 1. CT scan showing a large and expansive lesion in the head of the pancreas.
can be mistakenly diagnosed as a neuroendocrine tumor. Immunohistochemistry for light chains, associated with neuroendocrine markers, is essential for the differential diagnosis,\textsuperscript{10} and it was demonstrated in the present case.

EUS showed a hypoechoic, heterogeneous, well-defined mass in relation to the normal pancreatic echotexture, easily accessible to puncture, with the well-known advantages over CT-guided puncture. This is the first case described in the literature of successful EUS FNA in the diagnosis of plasmacytoma.

Treatment options include surgery, radiotherapy, chemotherapy, and combinations of them.\textsuperscript{11} Studies indicate that this tumor is highly radiosensitive and chemosensitive.\textsuperscript{2,12,13} However, as in our case and in other studies,\textsuperscript{11,14} the prognosis and the chemotherapy sensitivity can be poor and the disease course can be very aggressive.

References


\textsuperscript{7} Wilson TE, Korethkin M, Francis IR. Pancreatic plasmacytoma: CT findings. AJR 1989;152:1227-8.


