The association of bladder myeloid sarcoma and unclassified myelodysplastic/myeloproliferative disease

Mesanede myeloid sarkom ve sınıflandıramayan myeloproliferatif/myelodisplastik hastalık birlikteliği

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Abstract

Myeloid sarcoma of the urinary bladder is a rare disorder. We report a 71-year-old man with hematuria who had a diffuse myeloid sarcoma of the bladder. He was also under follow-up for unclassified myeloproliferative/myelodysplastic disorder, diagnosed two months before. Abdominal ultrasonography and computed tomography findings were normal. Diagnostic cystoscopy revealed patchy areas of mucosal swelling with hyperemia. Histopathological examination of biopsies demonstrated a neoplasm composed of blasts showing myeloperoxidase positivity by immunohistochemistry. To our knowledge, the current case is the first case of myeloid sarcoma in the urinary bladder without evidence of a mass lesion, with a concurrent diagnosis of unclassifiable myelodysplastic/myeloproliferative disease. (Turk J Hematol 2009; 26: 90-2)

Key words: Myeloid sarcoma, urinary bladder, unclassified myelodysplastic/myeloproliferative disease

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Özet


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Introduction

Myeloid sarcoma (MS) is a collection of myeloblasts or immature myeloid cells in extramedullary sites. The lesion may precede or occur concurrently with acute or chronic myeloid leukemia or the other types of myeloproliferative or myelodysplastic syndromes. The most common sites of involvement are soft tissue, orbit, lymph nodes and skin [1].

We present a rare case with both MS of the urinary bladder with patchy areas of submucosal involvement and an unclassified myelodysplastic/myeloproliferative disease.

Case Report

A 71-year-old man was admitted to the hospital with fatigue and weakness. Physical examination was unremarkable except for pallor. Laboratory parameters were as follows: hemoglobin (Hb) 7 g/dl, platelets 130x10^9/L and white blood cell (WBC) count 18x10^9/L with 70% neutrophils, 20% lymphocytes, 4% myelocytes, 2% normoblasts, 2% blasts, and 1% monocytes in the blood smear. Other laboratory investigations including urea, electrolytes and liver function tests were all in normal limits. Bone marrow aspiration and biopsy demonstrated marked myeloid hyperplasia with maturation and poor granulation of granulocytes, and normal number of blast cells (Figures 1, 2). Bcr/abl fusion gene was negative. The patient was accepted as unclassified myelodysplastic/myeloproliferative disease. The diagnosis of myelodysplastic/myeloproliferative disease, unclassifiable was based on bone marrow aspiration and biopsy examination (hypercellularity and marked myeloid hyperplasia with maturation), molecular study (bcr/abl gene fusion negativity), and laboratory and clinical findings (refractory anemia, leukocytosis, absence of monocytosis and hepatosplenomegaly, and no history of recent cytotoxic or growth factor therapy). No specific treatment was planned except blood transfusion when necessary.

Approximately two months later while under routine follow-up, the patient suffered hematuria. Anemia (Hb <8 g/dl) and leukocytosis (34x10^9/L) were still present. Abdominal ultrasonography and computed tomography (CT) of the abdomen were normal (Figure 3). Diagnostic cystoscopy revealed scattered erythematous mucosal hyperemia and swelling without any mass lesion. Biopsies were performed. Histological examination demonstrated a dense cellular infiltrate composed of immature cells under normal mucosal epithelium (Figure 4). The neoplastic cells were mostly medium in size with vesicular, round nuclei. The tumor cells were stained positive for myeloperoxidase (Figures 5, 6) and negative for cytokeratin, CD20, CD3, vimentin, and S-100 protein. No other foci of infiltration were detectable at the time of diagnosis or during follow-up. During follow-up, no other specific treatment modality in relation to hematological disease was planned due to the patient’s older age.

Discussion

Myeloid sarcoma is a collection of immature myeloid cells and was first described by Burns in 1811. MS was referred to as chloroma by King in 1853 due to its grossly green color when exposed to the air. This term was replaced with MS by Rappaport in 1966. In the new World Health Organization (WHO) classification, the term MS is used to define a tumor of myeloblasts or immature myeloid cells occurring in an extramedullary site [1].
Most MS cases occur in patients with a history of acute myeloid leukemia (AML), myeloproliferative disorder, or a myelodysplastic syndrome; however, it is rarely seen prior to the onset of bone marrow disease. The most common sites of involvement are soft tissue, lymph nodes, skin, periosteum and bone. Infrequently, it has been reported in other sites, such as the mediastinum, breast, orbit, uterus, epididymis, and gallbladder [2-4].

MS involving the urinary bladder is rare. In our review of the medical literature in English, we identified eight cases of MS involving the urinary bladder [2,3,5-9]. No cases were reported to have patchy submucosal involvement without any mass lesion, and there has been no reported association with myelodysplastic/myeloproliferative disease, unclassifiable. In the current case, abdominal CT demonstrated normal thickness of the urinary bladder wall and no mass lesion. Diagnostic cystoscopy demonstrated areas of patchy mucosal hyperemia.

MS is generally composed of poorly differentiated blasts and immature cells. The major differential diagnoses are large cell lymphoma, Burkitt lymphoma, lymphoblastic lymphoma and undifferentiated round cell tumor. Immunophenotyping by immunohistochemistry for expression of myeloid markers such as myeloperoxidase is essential for diagnosis of MS [10].

To our knowledge, this is the first report of a case with MS with diffuse infiltration of the urinary bladder and unclassified myelodysplastic/myeloproliferative disease. MS of the urinary bladder should be suspected in patients presenting with hematuria having myeloid leukemia and myeloproliferative or myelodysplastic syndromes, and even when associated with unclassified myelodysplastic/myeloproliferative disease, as in our case. Cystoscopic examination and biopsy of any suspicious mucosal lesion should be performed in the absence of any mass lesion in the urinary bladder in patients under follow-up for this diagnosis.

References