Immunoglobulin D plasmablastic myeloma mimicking metastatic malignancy

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Abstract

Immunoglobulin D (IgD) myeloma is an uncommon subtype of multiple myeloma and accounts for less than 2% of all myeloma patients. Plasmablastic myeloma is a rare morphological variant of multiple myeloma with a poor prognosis, and its association with IgD paraprotein is extremely rare. We report a patient who had the diagnosis of IgD myeloma with plasmablastic morphology and presented with multiple FDG-PET avid bone lesions, mimicking metastatic malignancy.

Case report

An 80-year-old lady was referred for anaemia and generalized bone pains. Blood investigations revealed low haemoglobin of 90 g/L, high calcium (2.72 mmol/L) and elevated creatinine (164 µmol/L). Eight years prior, the patient developed cancer in her left breast and attained remission after radical mastectomy and chemotherapy. Clinically there were no mass lesions in the contralateral breast, lymphadenopathy or abdominal masses. 18F-FDG-PET scan was performed due to the history of breast cancer, which revealed extensive FDG-avid uptakes throughout the skeleton, and in the bone marrow (Figure 1). Metastatic malignancy was the provisional diagnosis due to her past history of breast cancer. However no primary sites were identifiable and serum tumour markers such as CEA, CA19.9 and CA15.3 were within the reference ranges.

Since the differential diagnosis was multiple myeloma, serum protein electrophoresis was performed, which revealed a discrete paraprotein band in the gamma region. However, immunofixation electrophoresis was negative for standard heavy chain panel of IgG, IgM and IgA. But a monoclonal band was noted in the kappa light chain lane and Bence-Jones protein (kappa) was detected in urine electrophoresis. Hence IFE using anti-IgD and anti-IgE were performed and a discrete IgD paraprotein band was demonstrated and quantified as 8840 mg/l. There were no lytic or destructive bone lesions on skeletal CT scans despite multiple FDG-PET positive bone and marrow lesions. Bone marrow biopsy revealed diffuse infiltration by large and pleomorphic neoplastic cells containing eccentric nuclei, fine nuclear chromatin, prominent nucleoli and mitotic figures consistent with plasmablastic morphology (Figure 2). Immunohistochemical studies revealed strong surface positivity for CD138 on the neoplastic cells (Figure 3). IgD and kappa stains were also strongly positive in the cytoplasm of those cells (Figures 4 and 5), but negative for CD20, IgA, IgG, IgM and lambda and EBER. The diagnosis was plasmablastic myeloma of IgD-kappa subtype. Fluorescent in-situ hybridization testing on bone marrow was negative for high risk genetic rearrangements such as FGFR3-IGH, IGH-MAF and TP53. Serology tests for viral infections - HIV, Hepatitis B, Hepatitis C, EBV-IgM and CMV-IgM were negative. The patient was treated with bortezomib based regimen and bisphosphonates and she demonstrated good clinical response with drop in IgD paraprotein values to 1800 mg/l after 4 months; however, she developed fulminant septicemia and died two months later.

Discussion

Immunoglobulin D (IgD) myeloma accounts for less than 2% of all myeloma patients [1,2]. IgD myeloma is characterized by smaller size or absence of serum M-protein, predominance of lambda light chains, and high incidence of renal impairment, extraosseous involvement, hypercalcemia and amyloidosis [3-6]. Since conventional serum protein electrophoresis and immunoglobulin quantification assays are tailored to detect the more common IgG and IgA sub-types of myeloma, the diagnosis of IgD myeloma can be difficult [7].

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Figure 1. 18F-FDG-PET scan showed extensive FDG-avid uptakes in the skeleton
Plasmablastic myeloma is a rare morphological variant (~10%) of multiple myeloma and generally follows an aggressive clinical course with poor prognosis [8]. Morphologically plasmablasts differ from plasma cells due to the presence of larger nucleus size, fine nuclear chromatin pattern, prominent nucleolus, moderate rim of basophilic cytoplasm and absence of perinuclear hof [9].

In a study by Bartl et al., histological features of 674 cases of multiple myeloma were analyzed and classified based on cell size, cytoplasmic structure, and nuclear configuration of the neoplastic plasma cells. Plasmablastic type accounted for 2% and the rarest subtype in this series. This subtype showed the most aggressive clinical course among the morphological subtypes. Among plasmablastic myeloma, 60% had IgG paraprotein, 20% had IgA paraprotein, and the 20% with light chain, with no cases of IgD paraprotein IgD in the series [10].

Plasmablastic myeloma and plasmablastic lymphoma may exhibit similar morphological features histopathologically and cytologically with the neoplastic cells showing blastic features, with scanty plasmacytic differentiation [11]. Both these conditions demonstrate immunophenotype of plasma cells (CD38+, CD138+ and CD20-). However, plasmablastic lymphoma is usually associated with HIV infection and the malignant cells are EBV-infected in majority of cases and the main difference is EBV positivity in plasmablastic lymphoma, but negative in plasmablastic myeloma [10,12].

This case highlights the extremely rare association of IgD variant of multiple myeloma with plasmablastic morphology. Metastatic solid organ malignancy was initially suspected based on the presence of multiple FDG-positive skeletal lesions without lytic bone lesions, however further investigations confirmed IgD multiple myeloma.

**Author contributions**

Muhajir Mohamed - haematologist who diagnosed and treated the patient, performed literature review and wrote the manuscript. Wade Clarkson - biochemist who performed the diagnostic investigations, and checked the manuscript. Roya Arabi - clinical haematology registrar who helped in collecting details of the patient, and literature review and checked the manuscript.
Conflicts of interest

None to disclose for all authors.

References


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