

Anaplastic Multiple Myeloma- a Morphological Challenge – A Case Series

Abstract

Anaplastic multiple myeloma (AMM) is a rare and aggressive variant of plasma cell neoplasm. It is relatively treatment resistant when compared to conventional myeloma. It can arise *de novo* or transform from a preexisting plasma cell myeloma. Morphologically, AMM is recognized by the presence of immature plasma cells and pleomorphic, multilobated cells. Here, we present three cases of AMM.

Keywords: *Anaplastic myeloma, anaplastic variant of multiple myeloma, bizarre plasma cells, multilobated plasma cells*

**S Sakthisankari,
Prasanna N Kumar,
Tabbu Syed**

*Department of Pathology, PSG
Institute of Medical Sciences
and Research, Coimbatore,
Tamil Nadu, India*

Introduction

Anaplastic multiple myeloma (AMM) is a very rare morphological variant of multiple myeloma. It has an aggressive clinical course and poor prognosis. It is relatively more treatment resistant when compared to conventional myeloma. It can arise *de novo* or transform from a preexisting plasma cell myeloma.^[1,2]

Morphologically, AMM is recognized by the presence of immature plasma cells and pleomorphic, multilobated cells. These cells can be mistaken for carcinoma or lymphoma cells or even at times megakaryocytes. Here, we present three cases of AMM.^[3]

Case Reports

Case 1

An 85-year-old male who was diagnosed to have chronic kidney disease presented with the symptoms of anemia and back pain. Laboratory investigations showed anemia (hemoglobin [Hb]: 6.6 g/dl), erythrocyte sedimentation rate (ESR) of 120 mm, and elevated renal parameters (blood urea – 87 mg/dl and serum creatinine – 4.38 mg/dl). Serum albumin was 3.1 g/dl and globulin was 4.8 g/dl. Peripheral smear showed normocytic anemia with exaggerated rouleaux formation [Figure 1].

Bone marrow aspiration showed hypercellular marrow with significant

plasmacytosis (plasma cells – 72%). Also seen were many large cells with eccentrically placed hyperchromatic nuclei exhibiting marked nuclear lobulations and indentations. Subsequently, serum protein electrophoresis was done which demonstrated a discrete M band.

Case 2

A 61-year-old male, a diagnosed case of chronic kidney disease, presented with severe fatigue and bony tenderness. Blood investigations showed elevated renal parameters (blood urea – 95 mg/dl and serum creatinine – 7.42 mg/dl). There were significant albumin–globulin ratio reversal (serum albumin – 2.1 g/dl and globulin – 12.3 g/dl) and markedly elevated ESR (109 mm/h). Peripheral smear showed normocytic anemia (Hb – 4.4 g/dl), thrombocytopenia (87,000/mm³), white blood cell count of 4400/mm³, increased rouleaux formation, and presence of 4% atypical cells [Figure 2].

Serum protein electrophoresis showed a significant fall in albumin level and a steep increase in the gamma globulin region suggestive of “M band.” Bone marrow aspiration showed hypercellular marrow particles with pleomorphic plasmacytoid cells with lobated nuclei along with a few mature, binucleate and multinucleate plasma cells. Hypercellular marrow spaces with sheets and singly scattered pleomorphic cells were observed on trephine biopsy. Plasmacytoid morphology was focally discernible [Figure 3]. The differentials

Submitted: 13-Apr-2019

Revised: 24-Nov-2019

Accepted: 03-Jan-2020

Published: 27-Jun-2020

Address for correspondence:

*Dr. S Sakthisankari,
Department of Pathology,
PSG Institute of Medical
Sciences and Research,
Coimbatore - 641 004,
Tamil Nadu, India.
E-mail: sakthisankari@
gmail.com*

Access this article online

Website: www.ijmpo.org

DOI: 10.4103/ijmpo.ijmpo_96_19

Quick Response Code:



This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: WKHLRPMedknow_reprints@wolterskluwer.com

How to cite this article: Sakthisankari S, Kumar PN, Syed T. Anaplastic multiple myeloma- a morphological challenge – A case series. Indian J Med Paediatr Oncol 2020;41:430-3.

included lymphoma versus myeloma. Immunohistochemical markers were done. The cells expressed CD138 and were negative for CD20 immunostain [Figures 4 and 5].

Case 3

A 70-year-old male presented with severe fatigue, loss of appetite, and weight loss. He was a known case of multiple myeloma diagnosed 6 years back. He was in remission following treatment. Peripheral smear showed pancytopenia with increased rouleaux formation and presence of plasma cells. Bone marrow studies were done to rule out recurrence. Bone marrow aspirate showed bizarre plasma cells [Figures 6 and 7]. The marrow spaces showed diffuse infiltration by plasmacytoid cells and bizarre cells with multilobated nuclei. The cells expressed membranous staining for CD138. The bone marrow features were consistent with the anaplastic transformation of a myeloma.

Discussion

Multiple myeloma is a common hematological malignancy of the middle aged and elderly. AMM was originally used

to describe high-grade transformation occurring as a terminal event in the malignant plasma cell clones having pleomorphic morphology and immature plasma cells.^[3,4] It has been referred by various names such as plasma cell sarcoma, aggressive-phase myeloma, and dysplastic myeloma. It is more commonly reported in younger adults, with the median age being 53 years, associated with predisposition to extramedullary sites and immunoglobulin A isotype.^[2] Cases of anaplastic myeloma with poorly differentiated cells having extensive involvement of intra-abdominal and retroperitoneal sites have been reported in literature. AMM has morphological and flow cytometry features which are distinct from those of the conventional myeloma.^[4]

Historically, this disease has been very resistant to chemotherapy. It has been suggested that this aggressive phase could result from a clonal evolution of the original malignant cell and not considered as the development of an independent new neoplasia.

All the three cases reported were seen in adults. None of them had any evidence of extramedullary disease. Two

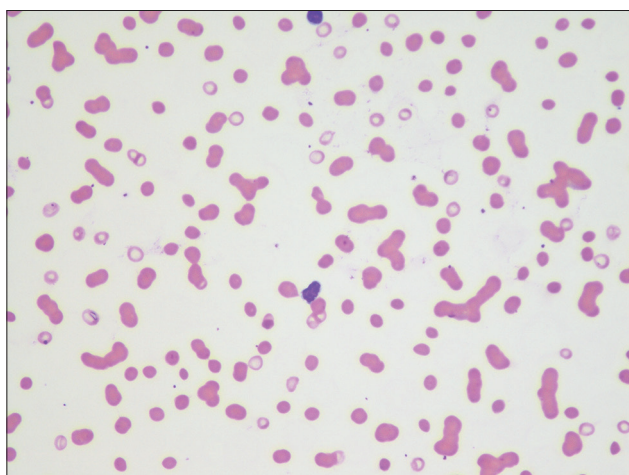


Figure 1: Case 1: Peripheral smear with exaggerated rouleaux formation ($\times 40$)

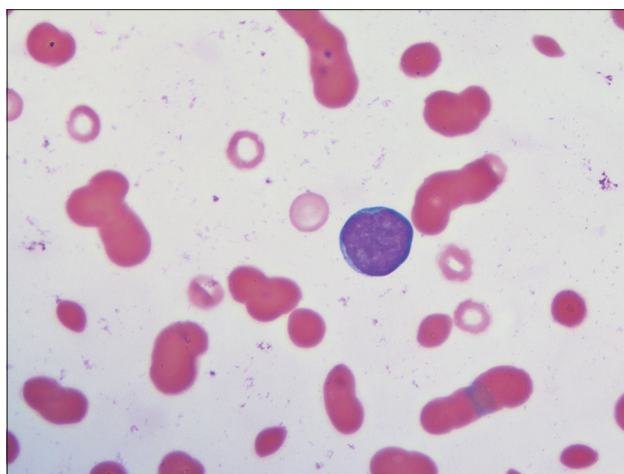


Figure 2: Case 2: Peripheral smear with a blast cell ($\times 100$)

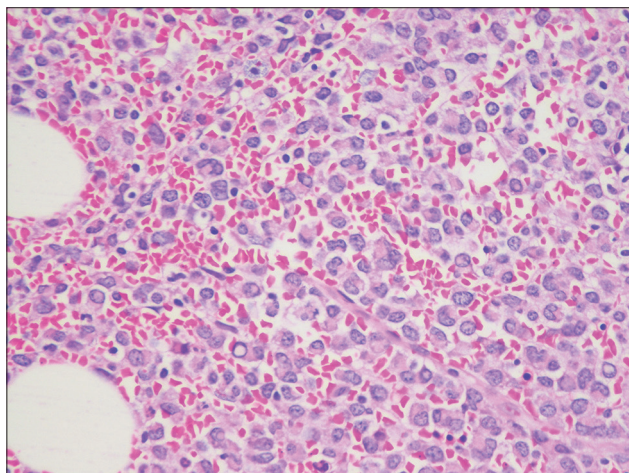


Figure 3: Case 2: Bone marrow trephine showing sheets of plasma cells, anaplastic cells, and Dutcher bodies (H and E, $\times 40$)

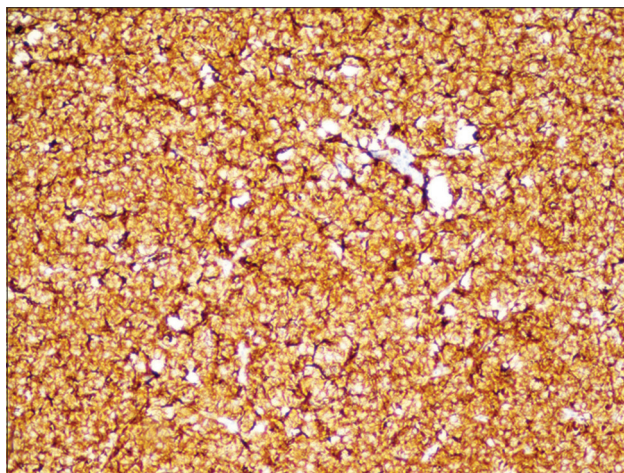


Figure 4: Case 2: Bone marrow trephine with cells expressing membranous CD138 (immunohistochemistry $\times 40$)

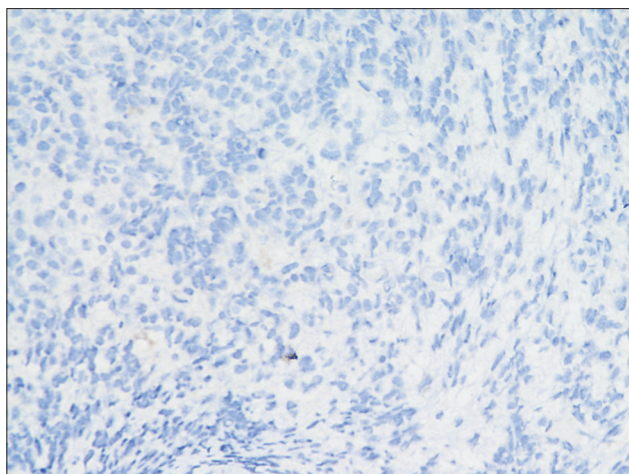


Figure 5: Case 2: Bone marrow trephine showing the neoplastic cells negative for CD20 (immunohistochemistry x40)

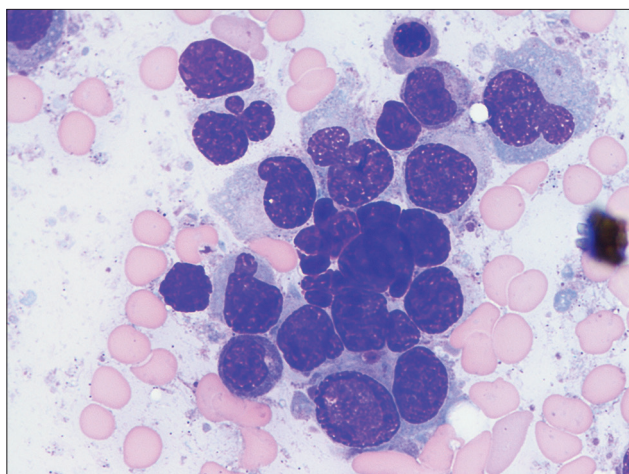


Figure 6: Case 3: Bone marrow aspirate with bizarre plasma cells (x100)

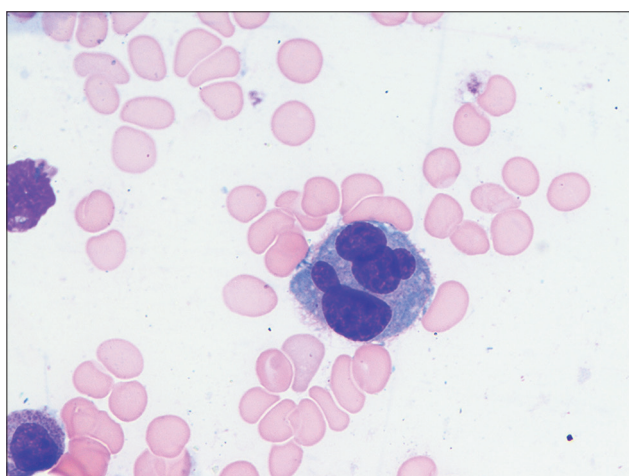


Figure 7: Case 3: Bone marrow aspirate with a plasma cell showing abnormal nuclear lobulation (x100)

cases presented *de novo*, whereas the other was anaplastic transformation seen in a patient in remission after treatment.

Multiple myeloma exhibits a broad morphological variation ranging from mature plasmacytic to highly anaplastic variants. The anaplastic variant poses a morphological diagnostic challenge as it mimics carcinoma and lymphomas, especially when presenting *de novo*. The presence of large cells with nuclear budding may mimic a megakaryocyte in bone marrow samples.^[5,6]

Case 2 reported here had a very anaplastic morphology, and the diagnosis was made with the aid of immunohistochemical staining with CD138.

Diagnosing a case as anaplastic myeloma without ancillary techniques is difficult, especially when the histology shows primitive cells. One of our cases had an undifferentiated morphology.

Bone marrow findings varied among the three cases.

Case 1 showed 72% plasma cells with the presence of few anaplastic cells. Whereas, case 2 showed predominantly large cells with pleomorphic morphology and scattered plasma cells.

Case 3 showed plasmacytoid cells with anaplastic morphology and occurred as a recurrence.

AMM, irrespective of presenting *de novo* or as progression, has been reported to have poor response to chemotherapy.^[4]

Conclusion

Here, we present a series of three cases of anaplastic myeloma for the rarity of this entity and also for the fact that these cases were considerably different from the previously reported cases in having no evidence of extramedullary disease and presenting at an older age. Anaplastic myeloma needs to be considered in bone marrow aspirates with pleomorphic morphology in a patient with other supportive evidence of plasma cell dyscrasia.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

Financial support and sponsorship

Nil.

Conflicts of interest

There are no conflicts of interest.

References

- Rodriguez C, Strowd RE, Grier DD. Anaplastic myeloma arising from myeloma: Successful management of two diseases in an individual. *Blood* 2017;130:5410.

2. Sethi S, Miller I. Plasma cell myeloma with anaplastic transformation. *Blood* 2016;128:2106.
3. Harankhedkar S, Gupta R, Rahman K. Pleomorphic multinucleated plasma cells simulating megakaryocytes in an anaplastic variant of myeloma. *Turk J Haematol* 2018;35:150-1.
4. Ammannagari N, Celotto K, Neppalli V, Lee K, Holstein SA. Anaplastic multiple myeloma: An aggressive variant with a poor response to novel therapies. *Clin Lymphoma Myeloma Leuk* 2016;16:e129-31.
5. Rao S, Kar R, Pati HP. Anaplastic myeloma: A morphologic diagnostic dilemma. *Indian J Hematol Blood Transfus* 2008;24:188-9.
6. Fujimi A, Nagamachi Y, Yamauchi N, Kanisawa Y. Morphological transformation of myeloma cells into multilobated plasma cell nuclei within 7 days in a case of secondary plasma cell leukemia that finally transformed as anaplastic myeloma. *Case Rep Hematol* 2017;2017:5758368.