

International Blood Research & Reviews

8(4): 1-3, 2018; Article no.IBRR.45424

ISSN: 2321-7219

Multiple Myeloma Relapse Presenting as Unilateral Blindness

Amit Badola^{1*}, Tanvi Khanna², Kunal² and Sanjiv Kumar Verma³

¹Department of Radiation Oncology, CRI, SRHU, Dehradun, Uttarakhand, India. ²Department of Pediatric Hematology Oncology and Stem Cell Transplant, CRI, SRHU, Dehradun, Uttarakhand, India.

³Department of Medical Oncology and Stem Cell Transplant, CRI, SRHU, Dehradun, Uttarakhand India.

Authors' contributions

This work was carried out in collaboration between all authors. All authors read and approved the final manuscript.

Article Information

DOI: 10.9734/IBRR/2018/45424

Editor(s):

(1) Dr. Mehmet Sonmez, Professor, Department of Haematology, School of Medicine, Karadeniz Technical University,

Turkey.

(1) Ilhami Berber, Malatya Education Research Hospital, Turkey.

(2) Carmino De Souza, Hematology and Blood Transfusion Center, University of Campinas, Brazil.

Complete Peer review History: http://www.sciencedomain.org/review-history/27535

Case Study

Received 19 September 2018 Accepted 23 November 2018 Published 01 December 2018

ABSTRACT

Multiple Myeloma is a chronic disease. While therapy is largely focused for control, relapse is inevitable. Central nervous system relapse of myeloma is less common. Unilateral painless blindness is a rare entity and its occurrence in the setting of multiple myeloma has not been commonly reported. We encountered a case of multiple myeloma on treatment, who developed unilateral blindness and later on confirmed to have relapse of disease.

Keywords: Multiple myeloma; unilateral blindness; central nervous system; extra-medullary relapse.

1. INTRODUCTION

Multiple Myeloma is a systemic malignant condition of plasma cell lineage. About 3% of

myeloma cases have extra medullary depositions of malignant plasma cells known as plasmacytomas [1]. While plasma cell deposits in upper half of skeletal system is common, the

*Corresponding author: E-mail: ammubadola@gmail.com;

involvement of central nervous system and orbits has been noted as well [2,3,4]. Unilateral optic nerve infiltration by plasma cells during treatment course of myeloma is an unusual and interesting finding.

2. CASE PRESENTATION

A 57 years old female presented with loss of vision from right eye for 2 months. She was a diagnosed case of multiple myeloma [IgG, Kappa] for last 2 years. Her initial treatment was with a 2 drug therapy including Bortezomib, Dexamethasone along with Zolandronic acid. She achieved remission after 4 months and was continued on same chemotherapy for total 8 months. She was placed on thalidomidedexamethasone regimen thereafter. She was on regular follow up and last assessment done 6 months back showed good control of disease. Her clinical examination now showed absence of light perception in right eye with no proptosis or ophthalmoplegia. Left ocular examination was normal. Detailed CNS examination showed no other significant finding. She underwent MRI which showed infiltrating compressing right optic nerve, distal to chaisma [Fig. 1]. Direct and indirect ophthalmoscopy was normal with normal fundus examination. Systemic workup showed increase in plasma cells in bone marrow and increased beta 2 microglobulin, serum LDH and IgG kappa chain. A systemic relapse of disease was considered along with plasma cell deposits over right optic nerve. She was started on Dexamethasone-Cyclophosphamide regimen in view of CNS penetration of these drugs. Radiotherapy to plasma cell deposit was started. Inj Bortezomib was added later in the course. She showed partial gain of vision post therapy. [Perception of light and finger counting at 1 meter present].

3. DISCUSSION

Multiple myeloma is a systemic disease of plasma cells. It is at one end of a spectrum of plasma cell dyscrasia, while milder localised forms can be detected as well. Being a systemic disease, its manifestation and presentation are variable. Usual presentation includes derangement of renal and haematological parameters as well as lytic bony lesions. However infiltration of organs has been noted as well [5,6]. It is practically an incurable disease and target of therapy is only to control it. Usual course with treatment is also with relapses at variable time periods. Ocular manifestation of multiple myeloma at presentation of relapse are variable. It includes corneal deposits, macular exudative detachments, or choroidal deposits. Usual presentation is with proptosis and hyperema [7,8]. Plasma cell infiltration of cranial

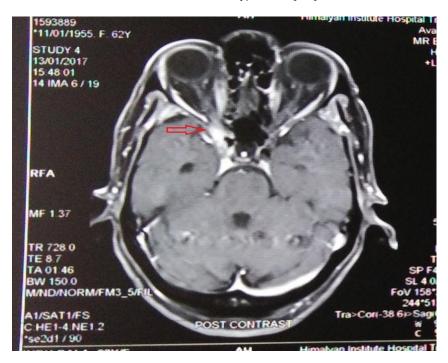


Fig. 1. MRI brain showing soft tissue density at right optic foramen

nerves has been noted less frequently. Frequent involvement is noted for oculomotor, abducens and hypoglossal cranial nerves [9,10]. Optic nerve compression causing blindness has been noted as initial feature in literature [11]. However a systemic relapse presenting with unilateral blindness is an unusual finding.

Treatment of CNS involvement is not standardised and mostly derived from systemic therapy of myeloma. Drugs with good CNS penetration should work well. As plasma cell deposits are sensitive to radiotherapy, an immediate relief is expected from palliative radiotherapy. Index case presented to us with unilateral blindness for 2 months duration. Post radiotherapy, only marginal improvement was noted. A possibility of long standing compression causing permanent damage to optic nerve fibres is considered.

4. CONCLUSION

Unilateral blindness due to malignant plasma cell deposits is an uncommon clinical entity. As multiple myeloma has vivid clinical presentation, a prompt suspicion should be kept for disease progression even in the scenario of unusual complaints.

CONSENT

As per international standard or university standard written participant consent has been collected and preserved by the authors.

ETHICAL APPROVAL

It is not applicable.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

 Adkins JW, Shields JA, Shields CL, Eagle RC Jr, Flanagan JC, Campanella PC.

- Plasmacytoma of the eye and orbit. Int Ophthalmol. 1996;20:339-43.
- Knapp AJ, Gartner S, Henkind P. Multiple myeloma and its ocular manifestations. Surv Ophthalmol. 1987;5:343-51.
- 3. Chim CS, Ng I, Trendell-Smith NJ, Liang R. Primary extramedullary plasmacytoma of the lacrimal gland. Leukemia Lymphoma. 2003;42:831-4.
- 4. Damaj G, Mohty M, Vey N, et al. Features of extramedullary and extraosseous multiple myeloma: A report of 19 patients from a single center. European Journal of Haematology. 2004;73:402–6.
- Ravinet A, Perbet S, Guièze R, et al. Lung postmortem autopsy revealing extramedullary involvement in multiple myeloma causing acute respiratory distress syndrome. Case Reports in Hematology. 2014;3. Article ID: 635237. DOI: 10.1155/2014/635237
- 6. Togano T, Sohtaro MI, Miwa A, Hagiwara S. Clinicopathological analysis of CNS involvement in multiple myeloma. Blood. 2015;126:5326.
- Malik A, Narang S, Handa U, Sood S. Multiple myeloma presenting as bilateral orbital proptosis. Indian J Ophthalmol. 2009:57:393-5.
- 8. Lazaridou MN, Micallef-Eynaud P, Hanna IT. Soft tissue plasmacytoma of the orbit as part of the spectrum of multiple myeloma. Orbit. 2007;26:315-8.
- Movsas TZ, Balcer LJ, Eggenberger ER, Hess JL, Galetta SL. Sixth nerve palsy as a presenting sign of intracranial plasmacytoma and multiple myeloma. J Neuro-Opthalmol. 2000;20:242-5.
- Kashyap R, Kumar R, Kumar S. Cranial nerve palsy in multiple myeloma and solitary plasmacytoma. Asia Pac J Clin Oncol. 2010;6:251-5.
- Yilmaz SG, Ture G, Zengin MÖ, Talay E, Men S. Optic nerve and dura mater involvement as the first sign of multiple myeloma. European Journal of Ophthalmology. 2014;25:77-9.

© 2018 Badola et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/4.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Peer-review history:
The peer review history for this paper can be accessed here:
http://www.sciencedomain.org/review-history/27535