



INTRAOURAL SUBLABIAL APPROACH FOR GIANT CELL REPARATIVE GRANULOMA OF THE LEFT ORBIT IN A 3 YEAR MALE CHILD.

Dr. Arsheed Hussain Hakeem	MBBS, MS, DNB, Department of Head and Neck Oncology, Apollo Cancer Institute, Hyderabad, India 500096
Dr. Hassaan Javaid	MBBS, Senior Resident, Department of Internal Medicine, Government Medical College, Srinagar, Jammu and Kashmir, India.
Novfa Iftikhar	MBBS, Department of Internal Medicine, Government Medical College, Srinagar, Jammu and Kashmir, India.
Dr. Usaamah Javaid	MBBS, Junior Resident, Department of Internal Medicine, Shadan Institute of Medical Sciences and research, Hyderabad, Telangana.

ABSTRACT We report a case of giant cell reparative granuloma of the orbit in a three year old male child and discuss its surgical approach. A 3-year-old male child presented to us with painless proptosis of the left eye. Magnetic resonance imaging demonstrated an intraosseous cystic orbital mass. Intraoral sublabial approach was used for complete excision of the lesion. At two years follow up, he has normal vision with no signs of recurrence.

KEYWORDS : Giant cell reparative granuloma; orbit; Intraoral; sublabial approach

INTRODUCTION

Giant cell reparative granuloma (GCRG) is a rare benign fibro-osseous lesion affecting facial bones especially mandible and maxilla.¹ Although, the aetiology of this rare lesion is far from clear, reactive response to intraosseous haemorrhage has been favoured as most likely cause.¹ On reviewing English language literature, only few reports describing its occurrence in the orbit can be seen.^{2,3} We report a case of GCRG arising in the left orbit causing painless proptosis in a 3-year-old male child. Intraoral sublabial approach was used to completely excise the lesion to avoid any scar or cosmetic deformity of the face. We also review the pertinent literature regarding management of this controversial lesion.

CASE REPORT

A 3-year-old male patient presented to our department of head and neck surgery with painless progressive proptosis of the left eye of three months duration (Figure 1). It was not associated with nasal discharge nor was there any history of trauma or significant paranasal sinus infections in the past. Clinical examination did not reveal any mass in the nasal cavity, or any palpable mass in the cheek substance. Laboratory studies including complete blood count, biochemical studies like serum calcium, parathyroid hormone, phosphate, alkaline phosphatase, were all within the normal limits. He underwent MR imaging studies of the orbit and paranasal. Coronal T1-weighted MR image demonstrates a heterogeneous mass involving left orbit with predominately low T1 signal intensity with internal septa (Figure 2).



Figure 1. Clinical Picture Showing Proptosis Left Eye.

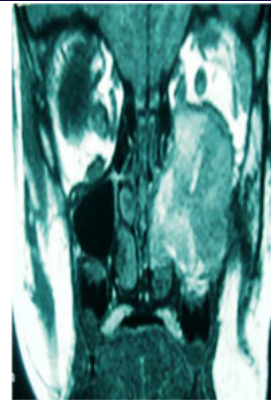


Figure 2. Coronal T1-weighted MR image demonstrates a heterogeneous but predominately low T1 signal intensity with internal septa involving left orbit.

Contrast-

enhanced sagittal T1-weighted MR image of left orbit demonstrates peripheral, septal, and soft tissue component enhancement on gadolinium administration (Figure 3). Orbital biopsy was performed which was consistent with GCRG. To completely excise the mass from the orbit, intraoral sublabial approach was used to access the mass to avoid any external scar. Histopathology of the specimen revealed a granulomatous lesion with reactive giant cells on a background stroma of plump spindle-shaped fibroblasts. The pathologic diagnosis was GCRG was made. The patient is free of recurrence at 3 years follow-up, and is being regularly followed in our clinic (Figure 4).

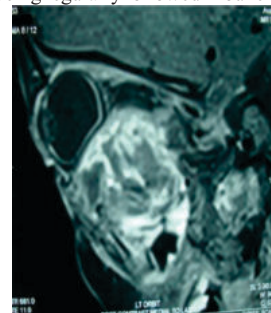


Figure 3. Contrast-enhanced sagittal T1-weighted MR image of left orbit demonstrates peripheral, septal, and soft tissue component enhancement.



Figure 4. Clinical Picture After 3 Years Of Follow Up.

DISCUSSION

Complete surgical excision rather than simple curettage is recommended treatment for GCRG, as curettage is associated with higher rate of recurrence.⁴ Surgical approach for GCRG affecting orbit in paediatric age group is of concern as it may result in deformity in the developing dentition or poorcosmesis. Several non-surgical methods have been proposed to treat GCRG, including radiotherapy,⁵ systemic calcitonin,⁶ intralesional corticosteroids injection⁷ and systemic α interferon.⁸ Among these nonsurgical treatment methods, intralesional corticosteroid injection has shown promising results and can lead to a complete resolution of the lesion or a significant reduction in size, allowing a more conservative surgery.⁷ We used intraoral sublabial approach for complete excision of the lesion, to avoid disfiguring scar on the face. Radiotherapy is contraindicated because of the potential for sarcomatous transformation that has been reported.⁹ Since there was not significant soft tissue extension and the lesion was close to the vital structures like orbit and optic nerve, we opted for surgical procedure and could completely remove the lesion. Now three years after the surgical resection patient has no symptom or signs to suggest recurrence. GCRG is a rare and benign fibro osseous lesion; therefore, definitive diagnosis will avoid mutilating radical surgery. Complete excision is preferred and should be done whenever possible as it decreases the chances of recurrence. For extensive or recurrent lesions involving surrounding soft tissues, treatment with intra-lesional injections of steroids can be useful in at least decreasing the size. More prospective controlled studies should be encouraged to find standard of care.

CONCLUSION

Intraoral sublabial approach is very useful in dealing with orbital and paranasal sinus lesion in pediatric age group. Not only it gives wide access, but also avoids external scar on face hence highly cosmetic.

Authors' Contributions AHH wrote the draft of the article. AHH, HJ, N and UJ helped in the final writing of the manuscript and gave final approval of the article.

All authors read and approved the final manuscript.

Compliance with Ethical Standards

Competing Interests: The authors declare that they have no competing interests.

Financial Support: None.

Consent: The informed consent was obtained from the patient for the publication of this report and any accompanying images.

REFERENCES

1. Jaffe HL. Giant cell reparative granuloma, traumatic bone cyst, and fibrous (fibro-osseous) dysplasia of the jaw bones. *Oral Surg Oral Med Oral Pathol.* 1953; 6:159–75.
2. Hoopes PC, Anderson RL, Blodi FC. Giant cell (reparative) granuloma of the orbit. *Ophthalmology* 1981; 88:1361-6.
3. Mercado GV, Shields CL, Gunduz K, Shields JA, Eagle RC Jr. Giant cell reparative granuloma of the orbit. *Am J Ophthalmol* 1999; 127:485-7.
4. Chuong R, Kaban LB, Kozakewich H, Perez-Atayde A. Central giant cell lesions of the jaws: a clinicopathologic study. *J Oral Maxillofac Surg.* 1986; 44:708–13.
5. Eisenbud L, Stern M, Rothberg M, Sachs SA. Central giant cell granuloma of the jaws: experiences in the management of thirty-seven cases. *J Oral Maxillofac Surg.* 1988; 46:376–84.

6. de Lange J, Rosenberg AJ, Van Den Akker HP, Koole R, Wids JJ, Van Den Berg H. Treatment of central giant cell granuloma of the jaw with calcitonin. *Int J Oral Maxillofac Surg.* 1999; 28:372–6.
7. Nogueira RL, Teixeira RC, Cavalcante RB, Ribeiro RA, Rabenhorst SH. Intralesional injection of triamcinolone hexacetonide as an alternative treatment for central giant-cell granuloma in 21 cases. *Int J Oral Maxillofac Surg.* 2010; 39:1204–10.
8. de Lange J, van Rijn RR, van den Berg H, van den Akker HP. Regression of central giant cell granuloma by a combination of imatinib and interferon: a case report. *Br J Oral Maxillofac Surg.* 2009; 47:59–61.
9. Adornato MC, Patcoff KA. Intralesional corticosteroid injection for treatment of central giant-cell granuloma. *Case report. JADA.* 2001; 132: 186-90.