Gaint Cell Granuloma: A Case Report

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Abstract

Central giant cell granuloma is a benign lesion which almost exclusively manifests in maxillofacial region. Usual site is anterior mandibular region. Radiographic features of this lesion closely resemble ameloblastoma and hyperparathyrodism. It is difficult to differentiate its histopathologic picture from that of hyperparathyrodism without co-relating clinically. This is a case report of central giant cell granuloma in body region of mandible along with its management and discussion (in literature).

Giant cell lesions of the jaw comprises of an interesting and controversial group.1 Among these giant cell granuloma is commonly encountered lesion. Any swelling which is not painful, gradually increasing in size and vascular in appearance raises the possibility of presence of giant cell granuloma. The case of peripheral as well as central giant cell granulomas are extensively well documented in literature2, and has been confirmed that 99% of giant cell tumor in jaws are giant cell granuloma. As it was presumed that it is an aftermath of trauma reparative word was frequently used (initially). In recent years term reparative has been deleted from original name given by Jaffe³ as it was found that lesion is basically destructive in nature rather than reparative.

Case Presentation

A 30 years old Indian female was referred to Oral & Maxillofacial Surgery Department for evaluation of swelling of size approximately 3.5x2.5 cm present in right mandibular region extending from right canine to 2nd molar, which was present since last 7 months and gradually increasing in size (Fig. 1).

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Fig: 1 (Extra oral pre-operative photograph)

On examination it was non-inflammatory in nature, hard in consistency involving both cortices. Swelling was localized, well defined, non tender with no associated paresthesia. First and second right molars were missing and premolars were slightly drifted. No other significant medical history was found except that patient was anaemic and seemingly undernourished.

Roentgenographie examination showed multilocular radiolucency with intervening septa's and well defined periphery (Fig. 2). FNAC suggested absence of any malignant change in lesion. Aspiration ruled out any cystic lesion.

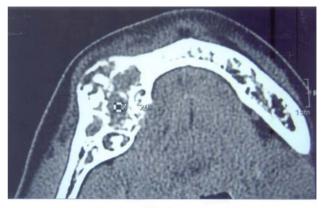


FIG: 2

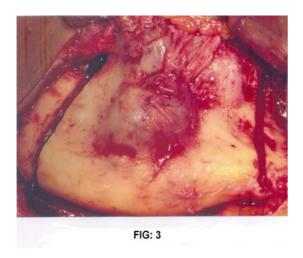
Axial section of CT scan of lesion

Differential diagnosis consist of (a) ameloblasgiant cell granuloma hyperparathyrodism (d) myxoma.

Possibility of hyperparathyrodism was ruled out by insignificant medical history, normal range of serum calcium and alkaline phosphtase level. Also, roentgenographically interlocular septa's were not straight (a characteristic of myxoma). More than usual bleeding during biopsy procedure increased suspicion of giant cell granuloma.

Histopathologic examination confirmed diagnosis of central giant cell granuloma with characteristic presentation of multinucleated giant cells, in loose fibrillar connective tissue and hemosiderin pigment foci present in between.

Treatment was done by segmental resection of involved part along with 1 cm of radiographically normal margin (Fig. 3,4). Frozen section from periphery was checked for absence of infiltrations. Primary reconstruction was done with the help of iliac crest grafting (Fig. 5,6).



Intra-operative (while resection)

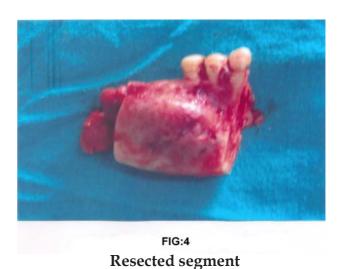






Fig: 6 (Post-operative extra-oral photograph)

malignant cell tumor and benign one includes fibrouso dysplasia ossifying fibroma, aneurismal bone cyst.4

Confusion regarding diagnosis was com-



Iliac crest in position (at continuity defect)

Discussion

In recent years there has been great controversy regarding different 1esions containing multinucleated giant cells. Variety of jaw lesions both malignant and benign contain multinucleated giant cells. Malignant one like osteogenic sarcoma, fibrosarcoma, fibrous histicytoma,

pounded by recognition of fact that hyperparathyrodism lesion histopathlogy is very similar to central giant cell granuloma. Distinction between giant cell tumor and giant cell granuloma is far from clear. Dron and Shafer concluded that they are probably same with one malignant variety. Bhaskar⁵ also felt that they are histologically indistinct, however Abrams and Shear⁶ said that there is significant morphologic difference between two. Lucas pointed out giant cell tumor have longer giant cells.

It is of general agreement that peripheral giant cell granuloma is much more common than central giant cell granuloma (5:1) and females are effected frequently (2:1) in comparison to men.

Mandible is commonly affected bone (2:1) with ante4rior segment most common to bear the brunt.⁷ Pain is not a prominent feature but bulging of cortices is present. Age is below 30.

Histopathology was described by Jaffe as giant cell lesion composed of fairly loose vascular stroma and small spindle shaped cells with hemorrhagic extravasation in between hemorrhagic extravastulation. Austin and Royer⁸ described bony stromal cells as elongated. I ueas remarked relative paucity and irregular distribution of giant cells in collagenous tissue.

Roentgenographically central giant cell granuloma is a destructive lesion with radiolucent area, ragged and sometimes smooth borders, sometimes showing fine trabeculae. Cortical plates are thin and expanded.

Traditionally treatment is done by curettage, eryosurgery ands excision. Hutler⁹ reported 67% cases treated by surgery alone and only 58% by curettage only. Bradley¹⁰ treated some of the lesions by conservative surgery and cryosurgery. Marconi¹¹ et al series included 25 cases of giant cell tumor treated with curettage plus eryosurgery using liquid nitrogen.

Currently central giant cell granulomas are

being treated successfully with curettage or surgery with isolated reports, where segmental resection with a margin of normal bone was done. John Webb¹² has described treatment of aggressive giant cell granuloma of mandible by combined curettage and eryosurgery.

Recurrence may be treated by repeat curettage. Percentage of recurrence is very low if surrounding area is cauterized.

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