Original article

A 38-year demographic study of central and peripheral giant cell granulomas of the jaws Saghafi S¹, Zare-Mahmoodabadi R², Ghazi N³, Zargari M⁴

Abstract:

Objective: The purpose of this study was to retrospectively analyze the demographic characteristics of patients with central giant cell granulomas (CGCGs) and peripheral giant cell granulomas (PGCGs) in Iranian population. Methods: The data were obtained from records of 1019 patients with CGCG and PGCG of the jaws referred to our department between 1972 and 2010. This 38-year retrospective study was based on existing data. Information regarding age distribution, gender, location of the lesion and clinical signs and symptoms was documented. Results: A total of 1019 patients were affected GCGLs including 435 CGCGs and 584 PGCGs during the study. The mean age was 28.91 ± 18.16. PGCGs and CGCGs had a peak of occurrence in the first and second decade of life respectively. A female predominance was shown in CGCG cases (57.70%), whereas PGCGs were more frequent in males (50.85%). Five hundred and ninety-eight cases of all giant cell lesions (58.7 %) occurred in the mandible. Posterior mandible was the most frequent site for both CGCG and PGCG cases. The second most common site for PGCG was posterior maxilla (21%), whereas anterior mandible was involved in CGCG (19.45%). The majority of patients were asymptomatic. Conclusions: In contrast to most of previous studies PGCGs occur more common in the first decade and also more frequently in male patients. Although the CGCGs share some histopathologic similarities with PGCGs, differences in demographic features may be observed in different populations which may help in the diagnosis and management of these lesions.

Keywords: central giant cell granuloma; peripheral giant cell granuloma

Bangladesh Journal of Medical Science Vol. 15 No. 02 April'16. Page: 220-223

Introduction:

Giant cell granuloma lesions (GCGLs) are benign, non-odontogenic, relatively uncommon tumors of the oral cavity, which arise either peripherally within gingiva, or centrally as an intraosseouslesion ¹. The peripheral giant cell granuloma (PGCG) is a rare reactive exophytic lesion arises in periodontal ligament and mucoperiosteum of the alveolar ridge. It is also known as a giant-cell epulis, giant-cell reparative granuloma ,or giant-cell hyperplasia². It occurs more frequent in the fifth and sixth decades of life with a slight female predilection³.

The central giant cell granuloma (CGCG) is a benign intraosseous proliferative lesion that occurs almost exclusively in the jaws. They comprise fewer than 7% of all benign tumors of the jaws ^{4,5}. This lesion mainly occurs in children or in youngadults, with a female predilection. It is more common in the mandible^{3, 6}.

CGCG and PGCG are virtually identical histologically, beingcharacterized by the presence of osteoclast-like giant cells scattered in a cellular fibrovascularstroma. However, despitetheir similarity, distinct clinical behavior

- 1. Shadi Saghafi, Oral and Maxillofacial Diseases Research Center, Department of Oral and Maxillofacial Pathology, School of Dentistry, Mashhad University of Medical Sciences, Mashhad, Iran
- 2. Reza Zare-Mahmoodabadi[,] Dental Research Center, Department of Oral and Maxillofacial Pathology, School of Dentistry, Mashhad University of Medical Sciences, Mashhad, Iran
- 3. Narges Ghazi, Oral and Maxillofacial Diseases Research Center, Department of Oral and Maxillofacial Pathology, School of Dentistry, Mashhad University of Medical Sciences, Mashhad, Iran
- 4. Mohammad Zargari, Private Office

<u>Corresponds to:</u> Narges Ghazi, Oral and Maxillofacial Diseases Research Center, Department of Oral and Maxillofacial Pathology, School of Dentistry, Mashhad University of Medical Sciences, Vakilalbad-Blv., P.O. Box 911735-984, Mashhad, Iran. **E-mail:** Ghazin@mums.ac.ir

is observed for these lesions.CGCGs are benign aggressivedestructive osteolytic lesionwith rapid growth, pain, root resorptionand tendency to recur after excision, whereas low recurrence rate and rare bone or tooth resorption are seen in PGCGs 7-9. There is considerable variation in the clinical behavior of CGCG. Rapid onset of pain, parasthesia, root resorption, and tooth displacement may be seen. Sometimes these lesions are asymptomatic 10. The distribution pattern of giant cell granulomas (GCGs) observed in one country may not be evident in other countries. The clinical and demographic features of these lesions including patient ageand sex, location of lesion, and distribution vary with race andgeographic location. There is little information in the English-language literatureabout the clinicopathologic features giant cell granuloma lesionsinIranian population.

The purpose of this study was to retrospectively analyzethe clinical features of 1019 patients with CGCGs and PGCGs in Iranian population. The findings were compared with the literature inrespect of age, gender, location of the lesion and clinical signs and symptoms (pain, swelling and bleeding).

Methods:

The data for the retrospective study were obtained from records of 1019 patients with CGCG and PGCG of the jaws referred to our department between 1972 and 2010. This 38-year retrospective study was based on existing data. Clinical data were analyzed, focusing on age, gender, location of the lesions, and clinical signs and symptoms includingpain, swelling and bleeding which were available in patients records. For GCG location the following scheme was used. The maxilla and mandible were divided into 6 anatomical regions, 3 on either side: anterior (from the midline to the distal surface of the canine), posterior (from the mesial aspect of the first premolar to the distal side of the third molar) and anterior-posterior. The anatomical region of 46 PGCGs and 60 CGCGs was not available. Statistical analysis was performed using SPSS software. Data were analyzed by applying γ 2 Test. Data were considered significant at P < 0.05.

Results:

A total of 1019 out of 9485 patients (10.7 %) were affected GCGLs during the study (435 CGCGs and 584 PGCGs). Patients ranged in age at the time of diagnosis from 2 to 90 years with a mean age of 28.91 ± 18.16 . PGCGs and CGCGs had a peak of occurrence in the first and second decade

of liferespectively. Table 1 shows distribution of PGCGs and CGCGs in different decades of agewith statistically significant difference (P < 0.05).

Table 1: Distribution of PGCGs and CGCGs in different decades of age

Age in decades	PGCG (n; relative %)	CGCG (n; relative %)
0-10	118 (20.20)	75(17.24)
10-20	97 (16.60)	111 (25.51)
20-30	90(15.41)	81 (18.62)
30-40	102 (17.46)	64 (14.71)
40-50	78 (61)	51 (11.72)
50-60	63 (13.35)	32 (7.35)
60-70	17 (2.91)	14 (3.21)
>70	19(3.25)	7(1.6)
Total	584(100)	435 (100)

A female predominance was shown in CGCG cases(57.70%), whereas PGCGs were more frequent in males (50.85%) with statistically significant difference (P < 0.05).

The distribution in terms of gender in CGCG and PGCG cases are presented in separately Table 2.

Table 2: Frequency of giant cell granuloma lesions based on sex origin

PGCG	Female (n; relative	287(49.15)
	%)	
	Male (n; relative %)	297(50.85)
	Total	584(100)
CGCG	Female (n; relative	251(57.70)
	%)	
	Male (n; relative %)	184 (42.30)
	Total	435(100)

Five hundred and ninety eight cases of all giant cell lesions (58.7 %) occurred in the mandible and 421 cases (41.3 %) were in maxilla (p<0.05). The mandibular and maxillary distribution of PGCGs and CGCGs are demonstrated in Table3.

Table 3: Frequency of giant cell granuloma lesions based on location

DCCC	M	224 (55 5)
PGCG	Mandible (n; relative	324 (55.5)
	%)	
	Maxilla (n; relative %)	260 (44.5)
	Total	584(100)
CGCG	Mandible (n; relative	274 (63)
	%)	
	Maxilla (n; relative %)	161 (37)
	Total	435 (100)
	111 1 1 0	• •

Posterior mandible was the most frequent site for both CGCG (35.75%) and PGCG (32.70%) cases.

Lesion	Mandible			Maxilla			
	Anterior	Posterior	Anterior-	Anterior	Posterior	Anterior-	Total
	(n;	(n;	Posterior(n;	(n;	(n;	Posterior(n;	
	relative	relative	relative %)	relative	relative	relative %)	
	%)	%)		%)	%)		
PGCG	98	176	28	99	113 (21)	24 (4.5)	538(100)
	(18.20)	(32.70)	(5.20)	(18.40)			
CGCG	73	134	22	65	59	22 (5.85)	375(100)
	(19.45)	(35.75)	(5.85)	(17.35)	(15.75)		_
P-valu	P=0.302			P=0.964			

Table 4: Anatomical regions of studied lesions in the jaws.

The second most common site for PGCG was posterior maxilla (21%), whereas anterior mandible was involved in CGCG (19.45%). PGCGs were distributed equally between the anterior maxilla and mandible. Table4 shows the frequency of studied lesions based on region of jawswithout statistically significant differences.

The majority of patients were asymptomatic. Bleeding was reported in 31% of PGCGs and 21% of CGCGs. Pain and swelling were only observed in 6.08% and 2.45% of patients respectively. Table 5 summarizes the clinical data of patients.

Table 5: Clinical data of patients

Lesion	Pain(n; relative %)	Swelling(n; relative %)	Bleeding(n; relative %)
PGCG	34(6)	9 (1.5)	180 (31)
CGCG	28 (6.45)	16 (3.65)	91 (21)
P-value	P=0.626	P<0.05	P<0.05

Discussion:

The present study details the profile of patients diagnosed as having central and peripheral giant cell granulomas between 1972 and2010. A total 1019 cases were evaluated, andepidemiologic findings were compared with previous studies. It is important to mention that we evaluated 1019 lesions in a 38-year period, whereas small number of cases considered in previous series^{1,6,9, 11-13}.

CGCGs occur more often in patients younger than 30 years of age^{9,14,15}. In our study, a peak of occurrence was in the second decade of life, corresponding to the findings of other authors¹¹. Although it has been shown that PGCGs occur more frequent in the fifth and sixth decades³the first decade was the most frequent age in the current study. In some studies patients were aged between four and seven decades, whereas most of our patients with PGCG were under 40 years old¹.

The majority of studies agree that there is a female predominance for CGCG lesions^{3,9,11,14,15}which is in agreement with our results. In the present study, PGCGs appeared more common in males, which is in contrast to the proved thesis that de-scribes predilection for female patients^{3,9,11,14-16}. Murat et al.'s also reported male predilection (56%) which was slightly higher than our results¹⁷.

A mandible predominance (58.7%) was identified in our series, and is in agreement with other studies^{4,9,14,17}.

Similar to our results, previous studies^{4,9,14,18}, have been stated that molar and premolarareas of mandible were more often affected by CGCGs than the anterior parts. Most of PGCG cases were also in the posterior part of mandible in our study. The second most common site for PGCG was posterior maxilla (21%), whereas anterior mandible was involved in CGCG (19.45%). In contrast to our results Boffano et al showed maxilla as the most frequent site for PGCGs¹⁹.

The clinical features of CGCGs varied considerably and is hard to predict⁹.

Bleeding was the most common clinical feature in our cases, whereas Sun et al reported asymmetric swelling of the jaw as the mostcommon clinical aspect in their series¹¹. Swelling was seen in only 3% of our cases (Table5).

Pain was considered to be associated with aggressive behavior of lesions(15)and was the second most frequent clinical aspect in our study. It should be mentioned that bleeding and pain were more common in patients with PGCG than CGCG.

In conclusion, in contrast to most of previous studies PGCGs occur more common in the first decade and also more frequently in male patients. Although the CGCGs share some histopathologic similarities with PGCGs, differences in demographic features

may be observed in different populations which may help in the diagnosis and management of these lesions.

Acknowledgment:

The research results given in this manuscript were

obtained from a doctoral thesis supported by the Vice Chancellor of Mashhad University of Medical Sciences, Iran.

Conflict of interest: None declared.

- 1. Etoz OA, Demirbas AE, Bulbul M, Akay E. The peripheral giant cell granuloma in edentulous patients: report of three unique cases. Eur J Dent2010; 4: 329-33.
- Chaparro-Avendaño Av, Berini-Aytés L, Gay Escoda C. Peripheral giant cell granuloma. A report of five cases and review of the literature. Med Oral Patol Oral Cir Bucal 2005; 10:48-57.
- Motamedi MH, Eshghyar N, Jafari SM, et al. Peripheral and centralgiantcellgranulomas of the jaws: a demographicstudy. Oral Surg Oral Med Oral Pathol Oral RadiolEndod2007;103:39-43.
- Kaffe I, Ardekian L, Taicher S, et al. Radiologic features of central giant cell granuloma of the jaws. Oral Surg Oral Med OralPathol Oral RadiolEndod 1996; 81: 720-6.
- Stavropoulos F, Katz J. Central giant cell granulomas: a systematic review of theradiographic characteristics with the addition of 20 new cases. DentomaxillofacRadiol2002; 31:213-7.
- 6. Nicolai G, Lorè B, Mariani G, et al. Central Giant Cell Granuloma of the Jaws. J CraniofacSurg2010; 21: 383-6.
- Kauzman A, Li SQ, Bradley G, et al. Central giant cell granuloma of the jaws: assessment of cell cycle proteins. J Oral Pathol Med 2004;33:170-6
- 8. Pandolfi PJ, Felefli S, Flaitz CM, et al. An aggressive peripheral giant cell granuloma in a child. J ClinPediatr Dent 1999; 23:353-5.
- Kruse-Lösler B, Diallo R, Gaertner C, et al. Central giant cell granuloma of the jaws: a clinical,radiologic, and histopathologic study of 26 cases. Oral Surg OralMed Oral Pathol Oral RadiolEndod2006;101:346-54.
- 10. Farrier SL, Farrier JN, Smart MK, et al. A 10-year review of the occurrence and treatment of central giant cell

- granulomas, in a District General Hospital. J Oral Pathol Med 2006; 35: 332-7.
- 11. Sun ZJ, Cai Y, Zwahlen RA, et al. Central giant cell granuloma of the jaws: clinical and radiological evaluation of 22 cases. Skeletal Radiol 2009; 38: 903-9.
- Kaplan I, Manor I, Yahalom R, et al.Central giant cell granuloma associated with central ossifying fibroma of the jaws: a clinicopathologic study. Oral Surg Oral Med Oral Pathol Oral RadiolEndod2007; 103:e35-41
- 13. Aghbali A, Sina M, VahidPakdel SM, et al. Correlation of histopathologic features with demographic, gross and radiographic findings in giant cell granulomas of the jaws. J Dent Res Dent Clin Dent Prospects 2013;7:225-9.
- De Lange J, Van den Akker HP. Clinical and radiological featuresof central giant-cell lesions of the jaw. Oral Surg Oral Med OralPathol Oral RadiolEndod 2005;99:464–70.
- Barnes L, Eveson JW, Reichart P, Sidransky D. World HealthOrganization classification of tumors: pathology and genetics ofhead and neck tumors. 3rd ed. Lyon: IARC; 2005.
- 16. AJ Mighell, PA Robinson, WJ Hume. Peripheral giant cell granuloma: a clinical study of 77 cases from 62 patients, and literature review. Oral Diseases 1995:1;12-19.
- 17. Murat H, Gongormus M, Abubekir H. Peripheral giant cellgranuloma; a clinical and radiological study. Pain Clin 2004;1:59-63.
- 18. Chuong R, Kaban LB, Kozakewich H, et al. Centralgiant cell lesions of the jaws: a clinicopathologic study. J Oral-Maxillofac Surg. 1986;44:708–13.
- 19. Boffano P, Benech R, Roccia F, et al. Review of peripheralgiantcellgranulomas. J CraniofacSurg2013;24:2206-8.