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# False gingival enlargement as a diagnostic problem: a case report

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Abstract: The aim of the case report was to describe gingival enlargement in a patient who came to the Department of Oral Medicine and Periodontology at Niš Dental Clinic. After anamnesis had been taken, and following clinical examination, laboratory blood analysis, radiological examination and pathological examination, it was established that gingival enlargement was a consequence of medicament injection. We are of the opinion that gingival enlargement was a consequence of sclerotic agent injection.

**Key words:** gingival enlargement; periodontal disease; sclerotic agents

### Introduction

Gingival enlargement is a common feature in gingival disease (1, 2). Many types of gingival enlargement can be classified in connection with aetiological factors and pathological changes (3-5).

- 1. Inflammatory enlargement: chronic and acute (6).
- **2.** Drug-induced enlargement (7–12).
- 3. Gingival enlargements associated with systemic diseases:
- (a) Conditioned enlargement (pregnancy, puberty, vitamin C deficiency, plasma cell gingivitis, non-specific conditioned enlargement-granuloma pyogenicum).
- **(b)** Systemic diseases causing gingival enlargement (leukaemia and granulomatous diseases - Wegener's granulomatosis, sarcoidosis, etc.) (13-15).
- 4. Neoplastic enlargement (gingival tumours) benign tumours and malignant tumours.
- 5. False enlargement these enlargements are not real enlargements, but may appear as such as a result of increase in size of

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© 2008 The Authors. Journal compilation © 2008 Blackwell Munksgaard the underlying osseous or dental tissue. The gingival tissue usually has no clinical signs of inflammation (16, 17).

Gingival enlargement is a fibrous overgrowth of gingival tissue that can be induced by various pharmacological agents through poorly understood mechanisms (9). It may occur because of hyperplasia or inflammatory processes. Unless a histological analysis is undertaken, the nature of the enlargement cannot be confirmed (6). Proliferative overgrowth of the gingival tissue makes it more difficult for patients to maintain oral hygiene (9, 18). Surgical correction of the gingival overgrowth is still the most frequent treatment. Such treatment is only advocated when the overgrowth is severe. It includes scalpel gingivectomy, overgrowth flap surgery, electro-surgery and laser excision (19).

# Case report

Patient ZG, male sex, 35 years of age, came to the Department of Oral medicine and Periodontology, Niš Dental Clinic because of the following problem: 6 months before, he had visited a private dentist in Leskovac with an intention of removing tooth calculus. After tooth calculus had been removed, the dentist injected a medicament into the gingival tissue once per week. After the third injection, the patient noticed that the gingival tissue started to enlarge and disturb him during toothbrushing.

Clinical examination: an abnormal bump, the size of a hazelnut, was present at the gingival area of the upper left second incisor and canines. The gingival tissue at this area was pale, firm and did not bleed at irritation (Fig. 1).

After removing the oral biofilm, obtaining laboratory results (which showed normal findings) and radiological examination, a surgery (removing of abnormal part of gingival tissue) was scheduled. It was noticed that the bone and periosteum were enlarged, and that they induced gingival 'swelling'. Taking into consideration the tumorous changes, a complete removal of the changed bone and gingival tissue from this area was conducted. The extirpated tissue was sent to histological verification. The histological analysis was performed by means of the standard haemotoxylin and eosin (HE) and it showed collagen enlargement in the gingival tissue and the presence of chronic inflammation (Fig. 2), as well as the bone structure which consisted of thin bone trabecules with osteoid at periphery. The post-operative follow-up was with no adverse events (Fig. 3).

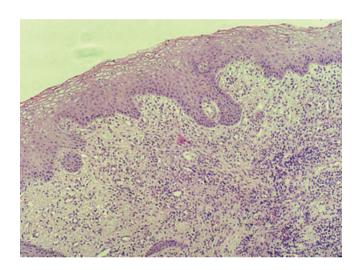


Fig. 2. Enlargement of collagen in gingival tissue. Focuses of chronical inflammation (HE, obj ×10).



Fig. 1. Clinical appearance.



Fig. 3. Clinical appearance after finished therapy.

# Discussion

Gingival enlargement can develop from chronic or acute inflammatory changes. Inflammatory gingival enlargement is usually secondary complication of other types of enlargement, and together they are called combined gingival enlargement. In these situations, it is very important to understand its double or multiple aetiology and to treat it adequately (3–5).

Acute inflammatory gingival enlargement results from bacteria which penetrated deeply into the tissue. Chronic inflammatory gingival enlargement is caused by prolonged exposure to oral biofilm and factors which are favourable for the accumulation of oral biofilm (20). Also, the administration of some drugs, such as anticonvulsants, immunosuppressant, calcium channel blockers, phenytoin, etc., can provoke gingival enlargement (18). Histological results of such enlargement show connective tissue and epithelium hyperplasia. In our case - the anamnesis, clinical investigation and histological analysis showed no reasons for such diagnosis. The histological analysis also showed bone structure which consisted of thin bone trabecules with osteoid at periphery. Such histological status points to the usage of sclerotic agents which injured periost, provoked subperiostal bleeding and later calcification.

Systemic diseases were excluded by diagnostic and laboratory tests (21). Tumours were excluded by histological investigation (11, 22-24).

The enlargement of the bone subjacent to the gingival area occurs most commonly in exostoses, but it can occur in Paget's disease, fibrous dysplasia, central giant cell granuloma, ameloblastoma, osteoma and osteosarcoma. In this case report, they were excluded by histological investigation, as well as radiological findings in this case report (4, 5, 11, 25-31). Because of the above-mentioned reasons, it can be assumed that the applied sclerotic agent therapy (the patient associates the time of gingival enlargement with the time of therapy) was the reason for gingival enlargement. It has been shown that a soft tissue injury in the vicinity of bone induces a periostal proliferation (30). Vascular disruption, as a consequence of a trauma, resulting in a transient ischaemia in the periosteum, would produce hypertrophy and hyperplasia of the periosteal cells, with an osteogen differentiation (28, 31).

Namely, classic periodontal therapy implies basic therapy with treatment of periodontal pockets. Also, other methods can be applied – surgical procedures, chemotherapeutics, antibiotics, physical methods, etc. (4, 5, 32-34). Some of these methods have had better and some worse results. But it is very important to have correct indications for certain method usage (18, 19).

Although the literature contains many references to phenytoin-, cyclosporine- and/or calcium channel blocker-induced gingival overgrowth, sclerotic agent-induced gingival overgrowth has not been reported. Similarly, an allergic reaction to a silicon-based dental material was noticed in a case report. It was shown that silicon can induce granulomatous reactions in the gingival tissue (35, 36). Deep application of sclerotic agents was first described by Hulin (cited by Petrović) (37). He described a range of sclerotic agents: calcium salts, especially phosphate salts, formaldehyde, hinine, urethane, methylene blue, etc. These agents were injected in the form of micro-drops using a sharp needle circularly around the tooth (in correlation with patient's anamnesis data). In this way, dense fibrous rings originate around the tooth. After retraction of the fibrous tissue, the depth of periodontal pocket reduces. Success cannot be achieved rapidly and improvements occur after few months, and both the patient and the therapist must have patience.

### Conclusion

If we want to avoid complications of periodontal therapy, it is necessary to have good knowledge of the anatomy-morphological characteristics of oral cavity, periodontal therapy techniques and possible unfavourable occurrences to which we have to react in a timely manner and to recognize the cause of gingival enlargement.

To the end of avoiding therapy mistakes, it is necessary to carry out diagnostic procedures and to avoid diagnostic problems such as problems caused by gingival enlargement.

## References

- 1 Beyer DJ, Belsito DV. Delayed hypersensitivity to silicon causing gingival hyperplasia. Contact Dermatitis 1997; 37: 234-235.
- 2 Armitage GC. Periodontal diagnoses and classification of periodontal diseases. Periodontol 2000 2004; 34: 9-21.
- 3 Seymour RA, Ellis JS, Thomason JM. Risk factors for drug-induced gingival overgrowth. J Clin Periodontol 2000; 27: 217-223.
- 4 Rees TD. Disorders affecting the periodontium. Periodontol 2000 1999; **21:** 145-149.
- 5 Somacarrera ML, Hernandez G, Acero J, Moskow BS. Factors related to the incidence and severity of cyclosporine-induced gingival overgrowth in transplant patients. A longitudinal study. J Periodontol 1994; 65: 671-675.
- 6 Majola MP, McFayden ML, Connolly C, Nair YP, Govender M, Laher MHE. Factors influencing phenytoin-induced gingival enlargement. J Clin Periodontol 2000; 27: 506-512.
- 7 Nishikawa S, Nagata T, Morisaki I, Oka T, Ishida H. Pathogenesis of drug-induced gingival overgrowth. A review of studies in the rat model. J Periodontol 1996; 67: 463-471.

- 8 Seymour RA. Effects of medications on the periodontal tissues in health and disease. Periodontol 2000 2006; 40: 120-129.
- 9 Brunet L, Miranda J, Farre M et al. Gingival enlargement induced by drugs. Drug Saf 1996; 15: 219-231.
- 10 Aimetti M, Romano F, Debernardi C. Effectiveness of periodontal therapy on the severity of cyclosporine A-induced gingival over growth. J Clin Peridontol 2005; 32: 846-850.
- 11 Embery G, Waddington RJ, Hall RC, Last KS. Connective tissue elements as diagnostic aids in periodontology. Periodontol 2000 2000: 24: 193-214.
- 12 Rees TD. Drugs and oral disorders. Periodontol 2000 1998; 18: 21 - 36.
- 13 Buckley DJ, Barrett AP, Bilous AM, Despas PJ. Wegener's granulomatosis - are gingival changes pathognomonic? J Oral Med 1987; **42:** 169-172.
- 14 Gonzales TS, Colema GC. Periodontal manifestations of collagen vascular disorders. Periodontol 2000 1999; 21:94-105.
- 15 Lee W, O'Donnell D. Severe gingival hyperplasia in a child with I-cell disease. Int J Paediatr Dent 2003; 13: 41-45.
- 16 Wright JM. Reactive, dysplastic and neoplastic conditions of periodontal ligament origin. Periodontol 2000 1999; 21: 7-15.
- 17 Khorsandian G, Lapointe HJ, Armstrong JEA, Wysocki GP. Idiopathic noncondylar hemimandibular hyperplasia. Int J Paediatr Dent 2001; 11: 298-303.
- 18 Prisant LM, Herman W. Calcium channel blocker induced gingival overgrowth. J Clin Hypertens 2002; 4: 310-311.
- 19 Mavrogiannis M, Ellis JS, Thomason JM, Seymour RA. The management of drug-induced gingival overgrowth. J Clin Periodontol 2006: 33: 434-439.
- 20 Shou S, Holmstrup P, Hjorting-Hansen E, Lang NP. Plaqueinduced marginal tissue reactions of osseointegrated oral implants: a review of the literature. Clin Oral Implants Res 1992; 3: 149-161.
- 21 Aarli JA, Tonder O. Effect of antiepileptic drugs on serum and salivary IgA. Scand J Immunol 1975; 4: 391-396.
- 22 Hancock RH, Swan RH. Nifedipine-induced gingival overgrowth. J Clin Periodontol 1992; 19: 12-15.

- 23 Hallmon WW, Rossmann JA. The role of drugs in the pathogenesis of gingival overgrowth. Periodontol 2000 1999; 21: 176-179.
- 24 Seymour RA, Smith DG, Rogers SR. The comparative effect of azathioprine and cyclosporine on some gingival health parameters of renal transplant patients. A longitudinal study. J Clin Periodontol 1987: **14:** 610-613.
- 25 Trackman PC, Kantarci A. Connective tissue metabolism and gingival overgrowth. Crit Rev Oral Biol Med 2004; 15(3): 165-175.
- 26 Elkhoury J, Cacchillo DA, Tatakis DN, Kalmar JR, Allen CM, Sedghizadeh PP. Undifferentiated malignant neoplasm involving the interdental gingiva: a case report. J Periodontol 2004; 75: 1295-1299.
- 27 Nitta H, Kameyama Y, Ishikawa I. Unusual gingival enlargement with rapidly progressive periodontitis. Report of case. J Periodontol 1993: **64:** 1008-1010.
- 28 Echeverria JJ, Montero M, Abad D, Gay C. Exostosis following a free gingival graft. J Clin Periodontol 2002; 29: 474-477.
- 29 McCarthy TL, Centrella M, Canalis E. Regulatory effects of insulin-like growth factors I and II on bone collagen synthesis in rat calvarial cultures. Endocrinology 1989; 124: 301-309.
- 30 Landry PS, Marino AA, Sadasivan KK, Albright JA. Effect of soft tissue trauma on the early periosteal response of bone to injury. J Trauma 2000; 48: 479-483.
- 31 Svindland AD, Nordsletten L, Reikeras O, Skjeldal S. Periosteal response to transient ischemia. Histological studies on the rat tibia. Acta Orthop Scand 1995; 66: 468-472.
- 32 Adrianes PA, Adrianes LM. Effects of nonsurgical periodontal therapy on hard and soft tissues. Periodontol 2000 2004; 3: 121-145.
- 33 Đajić D, Đukanović D, Zelić O, Ursu-Magdu I. Periodontal Disease. Gornji Milanovac, Dečje Novine, 1988.
- 34 Preus HR, Laurell L. Periodontal Diseases, a Manual of Diagnosis, Treatment and Maintenance. Chicago, Quintessence Books, 2003.
- 35 Beyer DJ, Belsito DV. Delayed hypersensitivity to silicon causing gingival hyperplasia. Contact Dermatitis 1997; 37: 234.
- 36 Btisch H. Silicone toxicology. Semin Arthritis Rheum 1994; 24:
- 37 Petrović L. Oral Diseases. Belgrade, Scientific Book, 1953.

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