Influence of Submandibulectomy on Alveolar Bone Loss in Rats

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Background: The incidence of dry mouth and its public health impact are increasing as the result of a progressively larger, medicated older population and because chronic diseases, like periodontitis, are prevalent pathologies among elderly patients. Periodontitis and continuous remodeling and rebuilding alveolar processes greatly affect the margin of the alveolar bone, and there is evidence indicating the role of submandibular glands in the regulation of immune/inflammatory reactions. The purpose of this study was to assess the effect of submandibular–sublingual complex ablation (Sx) on alveolar bone loss in rats submitted or not to ligature-induced experimental periodontal disease (EP).

Methods: Wistar male rats were submitted to Sx or sham operations (day 0). Two weeks later, unilateral EP was induced on the right mandibular first molars for 7 days with the contralateral side serving as control. Bone loss at the level of the dental pieces was estimated by bone histomorphometry on mesio-distally oriented sections of the molars and by the determination on lingual and vestibular mandibular surfaces of the distances from the cemento-enamel junction to the alveolar crest.

Results: Sx and EP significantly increased lingual and vestibular alveolar bone loss. Molars with EP exhibited greater lingual loss in Sx animals compared to those with the sham operation. EP induced similar interradicular bone loss in sham and Sx rats.

Conclusion: Sx has a deleterious effect on the periodontal tissues, particularly marginal alveolar bone, indicating the importance of the submandibular/sublingual glands in maintaining healthy periodontal conditions. J Periodontol 2008;79:1075-1080.

KEY WORDS
Bone and bones; periodontitis; saliva; salivary glands.

Saliva is an important protective factor for dentition and soft tissue because of its lubricating, cleansing, antibacterial, buffering, and remineralizing actions.1 The physiologic importance of saliva in oral health is clearly demonstrated in situations in which saliva production is impaired, such as in autoimmune diseases, e.g., Sjögren’s syndrome, or in patients receiving radiation treatment for head and neck cancer.2-5

Xerostomia or dry mouth is a condition frequently encountered in dental practices.6,7 The most common cause is the use of systemic medications that reduce saliva production.8,9 There is evidence that as the number of medications increases, a corresponding decrease in saliva output occurs, placing the elderly population, who are usually more medicated, at high risk for xerostomia.8,10,11 It is also well recognized that diminished salivary flow leads to difficulty in physiologic processes, such as mastication, swallowing, and speech,12 and can affect the quality of life as the result of problems related to the retention of dentures, infection with oral bacteria and yeast, development and progression of dental caries, and alterations in taste.2,13

Whole saliva is derived primarily from secretions of three pairs of major salivary glands, the parotid, submandibular, and sublingual glands, with a much smaller contribution from minor salivary glands and gingival crevicular fluid. Under resting conditions, which are prevalent in...
the oral cavity for most of the 24-hour period, submandibular glands are the major contributors to the volume of whole saliva (65%). Chronic xerostomia has a debilitating effect on the integrity of the hard and soft tissues of the mouth, and increasing evidence indicates that the submandibular glands exert regulatory influences on immune/inflammatory reactions. The spatial relationship between molars and the margin of the alveolar bone is greatly affected by the continuous remodeling and rebuilding processes in the alveolar bone and by periodontitis, an inflammatory periodontal disease that results in alveolar bone resorption and soft tissue attachment loss. To analyze the importance of submandibular/sublingual glands in maintaining healthy periodontal conditions, we assessed how the absence of submandibular endocrine and exocrine secretions, after submandibular–sublingual salivary gland complex ablation (Sx), affected periodontal bone loss in rats submitted or not to ligation-induced experimental periodontal disease (EP).

MATERIALS AND METHODS
Fifty-four adult male Wistar rats with an initial body weight of 230 to 250 g were randomly divided into two groups, experimental (Sx) and control (sham), and were maintained on standard chow pellets and tap water ad libitum. Housing conditions were as follows: galvanized wire cages; five animals per cage; temperature: 22°C to 24°C; humidity: 52% to 56%; and 12-hour light/dark cycles.

All experiments were performed following the National Institutes of Health guidelines for the care and use of laboratory animals (NIH 85-23, revised in 1985), and protocols were approved by the Ethical Commission of the Faculty of Dentistry, University of Buenos Aires.

Experimental Protocol
Rats were submitted to Sx or sham operations on day 0 (27 animals per group). Two weeks later, unilateral EP was induced on the right mandibular first molars of Sx and sham animals. The contralateral first molars (not ligated) of both groups (sham and Sx) were used as untreated controls. Animals were sacrificed by ether overdose 1 week later. Hemimandibles, once extracted and defleshed, were fixed in formalin for histomorphometry evaluation of interradicular bone (12 per group) or were stained with 1% aqueous methylene blue for microscopic determination of vestibular and lingual bone loss (15 per group).

Submandibulectomy
On day 0, the experimental group underwent bilateral Sx under light ether anesthesia. A midline incision of 15 mm was made through the skin and fascia of the ventral surface of the neck, the excretory ducts and main blood vessels were tied off, and the submandibular-

sublingual salivary complexes were dissected free from their supporting connective tissue and removed. They were excised with preservation of the surrounding neural structures, including the marginal mandibular branch of the facial nerve and the hypoglossal and facial nerves. The other group was subjected to a sham operation that involved full exposure of the glands and subsequent incision closure with a silk thread as in Sx animals.
**Periodontal Disease**
Two weeks after the onset of the experiment, animals were anesthetized with intraperitoneal administration of 2% xylazine hydrochloride (5 mg/kg body weight) and 5% ketamine hydrochloride (50 mg/kg body weight), and EP was induced by placing a cotton thread ligature around the neck of the lower right first molar. The cotton thread ligature was pushed into the gingival sulcus and was left in place for 7 days until sacrifice. The contralateral lower first molar without treatment (no ligature) was used as a control.

**Microscopic Examination of Periodontal Bone Loss: Distance Method**
Immediately after sacrifice, hemimandibles were resected, defleshed, and stained with 1% aqueous methylene blue to delineate the cemento-enamel junction (CEJ) and the alveolar crest (AC). A stereomicroscope** and a digital caliper†† were used to measure three lingual and three vestibular distances (mesial, central, and distal) from the CEJ to the most apical area of the AC (Fig. 1). Repeated determinations for this method yielded a relative coefficient of variation of 2%. The mean of the three recordings for each tooth surface was used as a measure of the total lingual and vestibular bone loss.

**Histomorphometric Evaluation**
Hemimandibles were fixed in formalin buffer immediately after resection. They were decalcified in 10% EDTA, pH 7, for 25 days. After this period, hemimandibles were dehydrated with ethanol and clarified with xylol. Finally, the sector containing the three lower molars of each decalcified hemimandible was embedded in paraffin at 56°C to 58°C. Under a stereomicroscope‡‡ and using a microtome§§ mesio-distally oriented sections of each lower first molar were obtained from paraffin blocks. Sections 7 μm in width were stained with hematoxylin and eosin, and histomorphometric evaluation was performed on digitized microphotographs using imaging software. The following static parameters were evaluated on the total interradicular bone: 1) bone volume (BV)/total volume (TV) (%) = fraction of TV corresponding to

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¶ König Laboratories, Buenos Aires, Argentina.
# Holliday-Scott, Buenos Aires, Argentina.
** Stemi DV4 Stereomicroscope, Carl Zeiss MicroImaging, Göttingen, Germany.
†† Digital caliper, Digimes, Geneva, Switzerland.
‡‡ Stemi DV4 Stereomicroscope, Carl Zeiss MicroImaging.
§§ Jung AG, Heidelberg, Germany.
|| Image Tool, University of Texas Health Science Center at San Antonio, San Antonio, TX.
bone tissue; total volume was taken as bone tissue plus bone marrow and periodontal ligament; 2) BV/TV (%) of section A (superior half of interradicular bone) (Fig. 2); and 3) height of the periodontal ligament (in micrometers); to measure the height of the periodontal ligament, four equidistant points were marked on the AC of the interradicular bone, and a line was drawn from each of the points to the bone (Fig. 2). The length of the lines was measured, and the mean value was calculated to obtain the height of the periodontal ligament of each section.19

Statistical Analysis
Results were analyzed statistically by the non-parametric Kruskal-Wallis and Dunn post hoc tests using a statistical program.¶¶ The level of significance was set at 0.05 and expressed as mean ± SEM (n).

RESULTS
The measures of lingual and vestibular bone loss, as distances between the CEJ and AC, revealed greater lingual than vestibular bone loss in sham and Sx animals (Figs. 3 and 4). In the unligated molars, greater bone loss was observed on lingual surfaces of Sx animals compared to sham animals (P <0.05; Fig. 4). The presence of the cotton thread for 7 days brought about a significant increase in lingual and vestibular bone loss (P<0.001). In addition, the lingual bone loss observed in EP molars of Sx animals was greater than in sham animals (P <0.05; Figs. 3 and 4).

Histomorphometric evaluation of interradicular bone revealed significant bone resorption in EP molars (P<0.05 and P<0.001 compared to the unligated control in sham and Sx animals, respectively) and no significant effect of Sx on molars with or without EP (Figs. 5 and 6).

DISCUSSION
The results suggest that the submandibular/sublingual glands are relevant in maintaining healthy periodontal conditions. Sx in rats increased lingual and vestibular bone loss as well as the lingual bone loss induced by EP, without affecting interradicular bone. The lingual alveolar bone loss was more pronounced than the vestibular bone loss in sham and Sx animals. Conversely, the percentage of bone loss induced by EP was greater on the vestibular surface. These results agree with the study of Ariji et al.,20 which demonstrated that the cortical plate changes induced by odontogenic infection were seen more frequently on the vestibular side than on the lingual side. The observation that molars with EP exhibited greater alveolar lingual loss in Sx animals than in sham animals suggests a local protective effect of the gland complex.

The submandibular gland produces serous, mucin-rich saliva that coats the oral tissues. Mucins represent the major organic component of submandibular/sublingual saliva. Their viscoelastic properties provide lubrication, and their high degree of glycosylation and potential for hydration prevent desiccation. As important components of the acquired pellicle and plaque matrix, mucins may also bind to toxins, agglutinate bacteria, and interact with host cells.21-23

A reduction in the output of mucins by Sx may increase mechanical forces and induce the loss of

¶¶ GraphPad Software, San Diego, CA.
supporting soft tissue, which, in turn, lead to loss of marginal alveolar bone as described herein.

Periodontitis is characterized by inflammation of the supporting tissues of the teeth and periodontal pocket formation that result in alveolar bone resorption and soft tissue attachment loss. In that sense, it has been suggested that the submandibular gland functions as a key regulatory organ in the oral neuro-immunoregulatory network. The submandibular gland produces, among others, nerve growth factor, epidermal growth factor, transforming growth factor-beta, and kallikreins, which are secreted into the saliva and the bloodstream and affect immune and mucosal tissue. Submandibular secretions enable the gland to exert regulatory influences on immune/inflammatory reactions. The submandibular/sublingual complex also produces antimicrobial proteins and secretory immunoglobulin A antibodies that are essential factors in the mucosal host defense. Suppression of all these secretions by Sx might alter the mucosal immune/inflammatory response and the regeneration and healing of periodontal support tissues, accounting for, at least in part, the negative effects of Sx on alveolar bone described herein.

In situations in which saliva production is impaired, such as in autoimmune diseases like Sjögren’s disease, a higher risk for periodontitis with higher plaque index scores, increased alveolar bone loss, and increased CEJ–AC distance have been reported. However, the occurrence of periodontal disease in Sjögren’s disease is still controversial. The health status of the gingival and periodontal tissues of these patients has also been described as similar to controls or with only slight differences. It is important to keep in mind that periodontitis is a multifactorial disease and, in most cases, a disease with a chronic progression. Exposure to factors that contribute to periodontitis occurs over a long period, so that, at the time of diagnosis, it may be difficult to identify and evaluate what cofactors have contributed to its development.

Our results indicated that the absence of submandibular secretions, exocrine and endocrine, have a direct deleterious effect on the periodontal tissues, which may increase the risk for gingival and periodontal pathologies. Such observations can have relevance, taking into consideration that the single most common side effect of many drugs is dry mouth and the evidence that as the number of medications per individual increases, there is a corresponding decrease in saliva output. Possibly, the increased incidence of diminished submandibular secretion in elderly patients favors that chronic diseases, such as periodontitis, are more prevalent. In this respect, the present observations of greater EP-induced lingual alveolar bone loss in Sx animals than in sham animals might be relevant.

No effect of Sx on periodontal disease was observed at the level of the interradicular bone; interradicular bone loss represents an advanced stage in the development of periodontitis in relation to marginal attachment bone loss. However, it is evident that rapid loss of the alveolar cortical plate induced by Sx is causal to progression to a more complex stage in the evolution of the disease.

How the removal of glands and their endocrine and endocrine secretions affect alveolar bone deserve further investigation. Sx might induce hyposalia, and/
or changes in salivary composition, and/or plaque accumulation on teeth, and/or alter the mucosal immune/inflammatory responses and/or calcium metabolism. Although the mechanisms remain unknown, our results emphasize the importance of submandibular saliva in maintaining a healthy oral environment and indicate that Sx has a deleterious effect on periodontal tissues, particularly marginal alveolar bone.

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