Early-diagnosed silent sinus syndrome and cone-beam computed tomography in a pediatric patient: a case report

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Chronic maxillary atelectasis (CMA) is a progressive alteration in the volume of the maxillary sinuses that may result in facial asymmetry. CMA in asymptomatic patients is known as silent sinus syndrome (SSS) and is a rare entity, especially in pediatric patients. This study reports a case of SSS in a pediatric patient who received an early diagnosis through cone-beam computed tomography (CBCT). An asymptomatic 12-year-old female patient in orthodontic treatment presented with opacification of the left maxillary sinus on a panoramic radiograph. Clinically, the patient had discrete hypoglobus and enophthalmos. CBCT and nasal video-endoscopy revealed ostiomeatal obstruction with bone deformity, leading to diagnosis of SSS. Endonasal endoscopic maxillary sinusotomy was performed. Two years later, the patient remained asymptomatic, and a second CBCT exam confirmed a stable condition. This case highlights the role of optimal radiographic interpretation for early diagnosis of maxillofacial alterations in pediatric patients.

Key words: Cone-beam computed tomography, Diagnosis, Maxillary sinus, Atelectasis, Surgery

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I. Introduction

The maxillary sinuses (MS) consist of a pair of pyramid-shaped cavities distributed bilaterally in the maxilla¹. The orbital floor, the alveolar process, and the lateral wall of the nasal cavity and facial surface of the maxilla represent the superior, inferior, and bilateral anatomic limits of the MS, respectively¹. Communication with the nasal cavity is through an ostium located high in the medial surfaces of these sinuses¹. Multiple functions of the MS are reported in the scientific literature, such as mucus production¹, improvement of voice quality², and participation in maxillofacial growth¹. Anatomic alterations in the MS, such as ostiomeatal obstruction, lead to functional deficits that have a negative impact on the internal pressure and aeration of the MS¹.

Radiographically, ostiomeatal obstruction may be observed in patients as opacification of the MS³. In routine dentistry, several circumstances may lead to sinus opacification³. However, in 1964, an unusual clinical entity with sinus opacification and enophthalmos was first reported⁴. Currently, this entity is known as chronic maxillary atelectasis (CMA) and manifests as a decrease in volume of the MS, progressing toward collapse of the sinus walls⁵. This alteration can be classified based on the following severity scale: stage I, membranous deformity; stage II, bone deformity; and stage III, clinical deformity⁶. CMA can be seen in both symptomatic and asymptomatic patients⁶. However, the latter is referred to as silent sinus syndrome (SSS), a subtype of CMA that combines facial asymmetry and lack of nasosinus symptoms⁶. Because patients with SSS are asymptomatic, the disease is often diagnosed late and culminates in inevitable facial asymmetry. Previous studies that explored the etiologies behind CMA reported chronic sinusitis, negative internal pressure, and valvular occlusion by the medial infundibular wall as potential causes⁵.
In dentistry, the MS are mainly examined with panoramic radiographs and cone-beam computed tomography (CBCT). These imaging modalities are routinely used in orthodontics, stomatology, maxillofacial surgery, and radiology for diagnosis, treatment planning, and follow-up. The detection and optimal interpretation of MS opacification on panoramic radiographs and CBCT scans are important in early diagnosis of SSS, particularly because patients are asymptomatic. Dentists must be aware of the radiographic and clinical signs of SSS to achieve early diagnosis. The present study reports a case of early-diagnosed SSS in an asymptomatic pediatric patient using CBCT.

II. Case Report

In November 2015, a 12-year-old female patient was referred to the local Department of Imaging for analysis of her left MS. Suspicion of a pathological alteration was raised after observing complete opacification of the left MS in a panoramic radiograph previously taken for orthodontic treatment. (Fig. 1)

During the anamnesis, the patient reported no symptoms, no systemic diseases, no previous medical history of therapeutic interventions or trauma in the MS, and no visual field defect. The extraoral physical exam revealed discrete midfacial asymmetry with hypoglobus and enophthalmos. (Fig. 2) The intraoral physical exam showed no infectious or inflammatory conditions in the teeth or soft tissues adjacent to the maxilla.

CBCT scans of the head were performed using an iCAT Next Generation (Imaging Science International, Hatfield, PA, USA) device with the following settings: field of view of 16×13 cm, voxel size of 0.25 mm, 37.07 mAs, 120 kVp, and image acquisition for 26.7 seconds. The images were analyzed by an otorhinolaryngologist and a maxillofacial radiologist using the iCat VisionQ (Imaging Science International) software package.

Bone deformity, ostiomeatal obstruction, and opacification of the left MS were observed on panoramic reconstruction of the CBCT scans (Fig. 3) as well as on axial and coronal slices. (Fig. 4) A nasal video-endoscopy performed in December 2015, confirmed the diagnosis of SSS. An endonasal endoscopic maxillary sinusotomy was performed under...
general anesthesia by an otorhinolaryngologist using 45° and 70° endoscopes to identify the main ostium and enable uncinectomy without injury to the periorbital bone. There were no transoperative complications, and postoperative analgesics and irrigation with saline solution were prescribed. Clearing of the ostiomeatal obstruction was observed and registered throughout 2-year follow-up. The patient remained asymptomatic, and no progression of facial asymmetry was detected during the follow-up period.

III. Discussion

SSS is a form of CMA that may be incidentally detected in patients with spontaneous midfacial eye asymmetry, such as enophthalmos or hypoglobus. Contrary to patients with CMA, those with SSS do not express nasosinus complaints. Diagnosing CMA or SSS is a challenging task. In fact, most patients are diagnosed late, when facial asymmetries are more evident. In this context, it is important to highlight the differential diagnoses of SSS in clinical practice. Clinically, SSS may be overlooked or interpreted as simple facial asymmetry, while radiographically, it may demonstrate signs similar to chronic maxillary sinusitis. The combination of clinical and radiologic evidence must be used to distinguish the disease; the exams must not be evaluated separately. In other words, although SSS may be clinically mistaken for facial asymmetry, the two entities can be distinguished radiographically with the aid of CBCT exams that may show incipient or late involvement of the MS. On the other hand, SSS may be radiographically mistaken for chronic maxillary sinusitis, but the two diseases can be distinguished based on typical clinical symptoms that are reported in chronic sinusitis but missing in SSS.

Attempting to explain the pathophysiologic process in SSS, the current scientific literature points toward hypoventilation of the MS secondary to obstruction of the ostiomeatal complex, directly affecting the process of gaseous exchange and leading to internal negative pressure. Next, damage in the bony limits of the MS, such as the floor of the orbit, can be induced by gradually increasing pressure, leading to further facial asymmetry. Routine dental imaging can reveal maxillofacial diseases, especially SSS because it is a progressive asymptomatic disease. The present study reported a case of SSS in a 12-year-old female patient who was examined for orthodontic treatment planning with the aid of a panoramic radiograph, allowing early diagnosis of SSS. A systematic review of 55 cases showed that the mean age of patients affected with SSS or CMA was 37.64±1.48 years.

Detecting alterations in the MS and diagnosing SSS are only possible with imaging. Several imaging modalities were used in the present case, namely panoramic radiograph (bi-dimensional modality), CBCT (three-dimensional modality),
and nasal video-endoscopy (non-radiographic modality). Each exam played an essential part in the diagnosis of SSS. Panoramic radiographs are consolidated as common imaging exams in routine dentistry. The present study highlights the importance of radiological interpretation in dental practice. Furthermore, analysis of radiographs must not be restricted to the teeth but also must include the maxillofacial bones. Proper knowledge on the interpretation of maxillofacial alterations is also essential. In the present study, the patient was referred to an otorhinolaryngologist and a maxillofacial radiologist who noted not only the radiographic alteration, but also the discrete enophthalmos and hypoglobus. These signs and the lack of symptoms, which had been previously reported in the scientific literature, led to suspicion of SSS. Thus, three-dimensional imaging was used for a more detailed analysis of the sinus morphology, and the inherent alteration of the left MS. CBCT revealed ostiomeatal obstruction and reduced limits of the left MS. Finally, video-endoscopy confirmed ostiomeatal obstruction finding and led to an endonasal endoscopic maxillary sinusotomy. Computed tomography and endoscopic surgery were previously used for diagnosis and treatment of similar maxillary alterations, which also resulted in positive outcomes.

Case reports of SSS are uncommon, especially in pediatric patients. Additionally, most of the cases in the literature were diagnosed in otorhinolaryngology or ophthalmology. The detection of maxillary alterations on panoramic radiographs is essential for early diagnosis of SSS. Dentists must be trained and aware of their role in prevention of severe sequelae that can result from late diagnosis of SSS. A parallel, multidisciplinary strategy from diagnosis to treatment of SSS must be considered for optimal outcomes.

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Consent for Publishing Photographs
Written informed consent was obtained from the patient for publication of this article and accompanying images.
Conflict of Interest

No potential conflict of interest relevant to this article was reported.

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