Up-to Date Review And Case Report

Characteristics of the odontological management of patients with progressive ossifying fibrodysplasia

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Abstract – Introduction: Progressive ossifying fibrodysplasia (POF) is a rare autosomal dominant disease characterized by the congenital malformation of the big toes and progressive postnatal heterotopic ossification of soft tissues with characteristic anatomical profiles. The maxillofacial region may also be affected. Observation: A 24-year-old man was referred by a traumatologist for the restoration of the oral cavity. He showed characteristic signs of POF. Oral clinical examination showed limitation of the oral opening, multiple dental caries, dental necrosis, and an asymptomatic impacted 48. Management included motivation for oral hygiene, scaling and dental extractions, followed by a prescription of steroidal anti-inflammatory drugs at a single dose of 2 mg/kg/day for 4 days. Discussion: The management of patients with POF in oral surgery has particularities. Dental care must be performed in brief sessions. The patient must be in a semi-sitting position, with the neck held upright, to avoid hyper extension of the neck and to improve comfort and safety. A prescription for corticosteroids is necessary after dental care to prevent possible heterotopic ossification. Through this article, we highlight the characteristics of POF, therapeutic attitude, and precautions to take to avoid possible complications.

Introduction

Progressive ossifying fibrodysplasia (POF) is a rare autosomal dominant disorder characterized by congenital malformation of the big toes and progressive postnatal heterotopic ossification of soft tissues with characteristic anatomic profiles [1–5]. The first heterotopic ossifications usually occur before the age of 10 years and mainly affect the tissues surrounding the hips, spine, and joints [6,7]. The outbreaks are episodic; immobility is cumulative. Heterotopic ossification usually begins at the age of 10 years. Temporomandibular joints (TMJ) can be affected early and are vulnerable to trauma at any age. The average lifespan of sufferers is 40 years; most deaths result from complications related to thoracic insufficiency syndrome or pneumonia via several mechanisms, such as costo-vertebral malformations, ossification of the intercostal muscles, or progressive deformation of the spine. Some deaths have been attributed to complications of general anesthesia due to intubation and ventilation difficulties. Currently, no medical or surgical intervention can alter the natural progression of POF [8].

The maxillofacial region can be equally affected. Trunical anesthesia, muscle fatigue due to extended periods during which the mouth is open can lead to progressive heterotopic ossification. The masticatory muscles (masseter, temporal, and buccinator muscles) as well as TMJ can be affected. Difficulty in opening the oral cavity and subsequent dietary difficulties may occur in addition to difficulties in maintaining oral hygiene [9,15,16]. Dental follow-up in patients with POF is essential because of the high risk of developing carious and/or periodontal lesions [10].

The objective of this article was to illustrate through a clinical case the characteristics of POF, therapeutic approach, and precautions to take to avoid possible complications.

Observation

The case was a young man aged 24 years who consulted for issues with his oral cavity. His illness was discovered at the age of 1 year following a cervical trauma. The patient underwent two hip surgeries in 2010 and 2016, following alterations of the hip and hip joint mobility.

He presented characteristic clinical signs of POF: decreased mobility of the peripheral and spinal joints, laterodeviation of the hallux (Fig. 1), and deformities following heterotopic ossifications in his back (Fig. 2) and hand (Fig. 3).
Oral clinical examination revealed a restricted jaw opening (20 mm), carious lesions on 11-12-23-27-36-37-38-46, pulpal necrosis on 17-14-35-45, and an asymptomatic impacted 48 (Figs. 4 and 5).

Radiological examination revealed two well-defined radiographic images related to 35 and 45 (Fig. 6).

The proposed treatment comprised improved oral hygiene, descaling, extractions of 14-35 and 45, endodontic care of 23 and 17, and restorative care of 11-12-36-37-38-46 using glass ionomer cement.

Fig. 1. Laterodeviation of the hallux.

Fig. 2. Deformations related to heterotopic ossifications in the back.

Fig. 3. Heterotopic ossifications in the hand.

Fig. 4. Intraoral view of the upper jaw.
The surgery selected was the least traumatic possible in the given circumstances. The patient was placed in a semi-sitting position, with the neck held straight to prevent hyperextension of the neck. The appointments were kept short. After descaling and extractions, a prescription of steroidal anti-inflammatory drugs (prednisone) was made at a single dose of 2 mg/kg/day for 4 days.

A 6-month follow-up did not reveal any heterotopic ossifications resulting from the procedures.

Discussion

POF is a rare genetic disease that affects approximately one in two million people. This disease is characterized by skeletal malformations and progressive heterotopic ossification of the skeletal muscles, tendons, ligaments, and fascia \[11,12\]. There is no ethnic, racial, sexual, or geographical predisposition \[13\]. POF is caused by genetic mutations inducing osteogenic differentiation \[14\]. Two clinical features define classic POF: big toe malformations and progressive heterotopic endochondral ossifications in key anatomical areas (back, knees, and hands). In our case, the patient presented characteristic signs of POF decreased mobility of the peripheral joints and the spine, hallux laterodeviation, and heterotopic ossifications in the hand and back.

Individuals with POF show no signs at birth, with the exception of big toe malformations that are present in all affected individuals. During the first 10 years of life, most affected children develop episodic and painful swellings of the soft tissues \[15,16\]. In our patient, the diagnosis was made at the age of 1 year, following heterotopic ossification at the cervical level after a trauma.

While some outbreaks spontaneously regress, most transform soft connective tissue, including fascia, ligaments, tendons, and skeletal muscles into mature heterotopic bone. Heterotopic bone replaces skeletal muscle and connective tissue with via endochondral ossification, leading to bone encapsulation and permanent immobility \[15,16\]. Individuals with POF experience dental problems similar to the general population. However, if there is a limitation in the movement of the jaw or ankylosis of TMJ, these patients may have difficulty with brushing and maintaining oral hygiene; therefore, the risk of carious and/or periodontal disease increases in these patients \[10\]. Restorative dental care is technically difficult to perform. The use of a filler material that releases fluoride is recommended. It binds to the tooth and releases fluoride into the surrounding tooth structure, preventing new tooth decay \[16\].

In our case, the patient showed restricted jaw movement, explaining the presence of multiple caries and pulpal necrosis due to difficulty in brushing and maintaining oral hygiene. We proposed ionomer glass fillings to limit the risk of carious recurrence. Minor trauma can trigger new painful outbreaks of POF, leading to progressive heterotopic ossification \[15,16\]. Therefore, the intramuscular injection of a local anesthetic is to be avoided because it is extremely toxic to skeletal muscles. Truncular anesthesia at the Spix spine can therefore lead to heterotopic ossification \[17\]. Infiltration anesthesia is difficult to administer in posterior mandibular regions of permanent teeth, but an intra-ligamentous anesthesia may be useful. The use of general anesthesia has been described, but deformities of the spine and involvement of the respiratory system seriously contraindicate or complicate its use \[18\]. We performed dental extractions in the least traumatic way possible with apical anesthesia.

Permanent ankylosis of the maxillae may occur following minimal soft tissue trauma during routine dental care or due to a poorly adapted position \[17\]. During dental care, short appointments should be scheduled, the patient should be as comfortable as possible, and the chair should be positioned at 45° to avoid excessive neck and lower jaw extension and therefore prevent possible muscle exertion \[19\]. In our case, following this procedure, we did not find ossification in any of the at risk sites.

Steroidal anti-inflammatory drugs are indicated as the first-line therapy immediately following any inflammatory episodes. High-dose corticosteroids (2 mg/kg/day prednisone equivalent) \[20\], initiated within 24 hours after an active flare-up, for a duration of 4 days may help reduce intense inflammation and tissue edema in the early stages of the disease. The use of
corticosteroids should be limited to the treatment of relapses that affect the major joints, jaw, or sub-maxillary region. In our patient, we followed the same prescription protocol in a preventive way without any problems with cicatrization or heterotopic ossifications.

Other authors advocate that the use of nonsteroidal anti-inflammatory drugs (NSAIDs) in symptomatic cases during the episodes. The use of mast cell inhibitors and aminobiphosphonates in this context is possible [20].

Conclusion

Preventive oral measures are essential in patients with POF, specifically during childhood. Regular brushing is essential. The use of toothpaste, gel, fluoride-based mouthwash, and fluoridated water is highly recommended in these patients [19]. Educating patients with POF as well as their parents and educators regarding preventive oral measures is paramount to establish good oral hygiene and facilitate their care.

Conflicts of interests: The authors declare that they have no conflicts of interest in relation to the publication of this article.

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