Fibrous Dysplasia of the Middle Turbinate: About A Case Report
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Fibrous dysplasia is a common benign fibro-osseous disease involving the flat bones, often affecting the bony structures of the skull and facial skeleton. Primary occurrence or secondary involvement of the nasal turbinate is not a common manifestation of the disease. Involvement of the inferior turbinate generally does not have specific management-related issues; however, involvement of the middle turbinate, especially the lateral lamella, can predispose to surgical morbidity during endoscopic surgical management. Clinical presentation, management and features of the disease on CT imaging are presented.

Keywords: Fibrous dysplasia, CT, MRI.

INTRODUCTION

Fibrous dysplasia (FD) is a condition characterized by progressive replacement of the normal bony structures with benign cellular fibrous connective tissue, leading to disorganized structure of the bone.1 Depending on the extent of involvement of the skeletal components, the disease can be categorized as monostotic (limited to a single bone) or polyostotic[1, 3]. The most common sites of involvement are the membranous bones of the pelvic girdle, femur and tibia. Skull is also a common site of involvement. Involvement of the nasal turbinate is infrequent [3, 5]. Isolated involvement of the inferior turbinate, in view of its anatomical relation, does not predispose to significant management problems. However, involvement of the middle and superior turbinate, and lateral and basal lamella creates a unique situation wherein there is involvement of the skull base structures, leading to increased surgical morbidity. Involvement of the lateral lamella, by virtue of its attachment to the cribriform plate and proximity to the ethmoidal arteries, predisposes to risk of damage to the floor of the anterior fossa and vascular injury during surgical or endoscopic intervention. We present the case of fibrous dysplasia of the middle turbinate in a 11 year old child.

CASE REPORT

- A 11-year-old male was admitted in our radiology department with difficulty in breathing through the nose for 2 years.
- Physical examination found a hard mass protruding through the right nostril.
- Laboratory parameters, including levels of alkaline phos-phatase, were normal.
- The CT showed ( figure 1):
  - a voluminous ground-glass bony lesion with frosted glass on the right nasal cavity which i
  - visualization of the turbinate.
- It is responsible for an ostiomial filling with fluid retention of the right maxillary sinus without visualization of the middle turbinate.
- This mass is well limited measuring 34x52x 47mm.
- She pushes the nasal septum to the left with no extension beyond the skull-base or into the orbit.
- The patient underwent endoscopic surgery under general anaesthesia and complete excision of the mass was performed
- The mass was adherent to the medial wall of the orbit, the posterior aspect of the septum and the medial surface of the inferior turbinate.
- Histopathological examination confirmed the diagnosis (figure 2).
DISCUSSION

Localization of FD only at the middle turbinate is an extremely rare event; only two other cases having been reported in the English literature[2, 6]. Neither of which underwent surgical treatment. Nevertheless, the incidence of this disease is probably underestimated due to the asymptomatic bony proliferation that remains undetected until it gives rise to pathological fractures, pain due to neurological compression or functional disorders in the affected organ. Therefore, without related symptoms, diagnosis is usually incidental during radiological examinations performed for other causes.

On the middle turbinate, the bony degeneration gives rise to an increase in volume that can cause nasal obstruction or alterations in sinus drainage. Videorhinoscopy shows turbinate enlargement that can be mistaken for a concha bullosa; this examination also allows evaluation of other, associated or correlated, sinus-nasal diseases. Axial-coronal CT, in thin sections, is the radiological exam of choice: it highlights the bony nature of the lesion with the characteristic “ground-glass” appearance. The radiographic findings may change in relation to the evolutive stage of the disease, therefore, three classic radiological patterns have been described:
- Lytic or cysti-like form characterized by roundish rarefactive areas with a sclerotic border;
- Sclerotic form with homogeneous bone expansion;
- Pagetoid form with dishomogeneous areas of radiodensity and radiolucency [1, 4].
- With CT, it is also possible:
  - to define the boundaries and connections of the lesion;
  - to exclude a polyostotic craniofacial form;
to identify other possible associated disorders;

to monitor evolution of the disease.

Classically, on MRI, FD shows hypointensity in T1 and T2 weighted sequences, owing to a large fibrous component of the lesion. However, signal intensity of the bone lesions can vary depending on the composition of the bone and extent of mineralization.

FD has to be differentiated from other fibrodysplastic disorders (Paget disease, hyperparathyroidism, osteogenesis imperfecta …) and from ossifying lesions (osteoma, osteosarcoma, chondroma, ossifying fibroma …). To this end, histopathological studies are of fundamental importance since they reveal the presence of irregular bony trabeculae embedded in a cellular fibrous stroma without osteoblastic rimming. However, to distinguish diseases with such similar clinical and histological features, close cooperation between the clinician, the pathologist and the radiologist is essential. Moreover, especially for the ossifying fibroma that, unlike FD, usually occurs in adult age, is characterized by monostotic expansive lesions with a well-defined edge and, for this reason, shows a better response to surgical treatment.

FD may be associated with hormonal changes as well as defects in the calcium-phosphorous metabolism; therefore, blood tests such as calcemia, alkaline phosphatase and parathormone assessment are mandatory to exclude a syndromic form, in particular those of McCune-Albright. On the contrary, the association of FD with polyposis sinusitis, allergic bronchial asthma and ASA intolerance (Widal triad), that we consider incidental, has not been described, so far, in the literature.

No medical treatment has been found to be effective in the management of FD; therefore, in those cases presenting localized lesions with functional disorders or cosmetic alterations, surgical treatment is indicated. On account of the poorly defined limits between normal and dysplastic bone, “en-bloc” excision is often very difficult and associated with recurrence in approximately 20-25% of these patients. Partial resections or bony reshapes are more frequently performed also in consideration of the fact that the disease tends to exhaust itself after puberty.

Surgical treatment, particularly if not radical, requires long-term follow-up in order to monitor the risk of recurrence and to evaluate possible malignant degeneration. Indeed, FD is associated with sarcomatous evolution in 0.5-1% of cases, in particular in cases of craniofacial lesions [9, 10]. The risk of mali nant degeneration increases considerably after radiation-therapy that must be avoided in all cases.

In conclusion, even though exceptional, localization of FD at the middle turbinate has been described, and, therefore, must be borne in mind in the differential diagnosis of the craniofacial ossifying disorder 2-4. Likewise, possible syndromic associations should be sought.

Videorhinosenoscopy and craniofacial CT are fundamental examinations in the correct assessment of this disease, diagnosis of which, however, depends on a scrupulous study of the histopathological features.

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