Fibrous dysplasia of the right maxillary bone

Tamer W Kassem
Department of Diagnostic and Interventional Radiology, University of Cairo, Egypt

Abstract

Fibrous dysplasia is a rare bone disease caused by mutations in GNAS gene. The normal bone is replaced with fibrous tissue and abnormal (woven) bone. Monostotic disease occurs in one bone while polyostotic disease occurs in multiple bones [1]. It is most common in ribs (28%) and occurs in craniofacial bones in 10 to 25 % of cases [2]. Craniofacial fibrous dysplasia (CFD) lesions are detected early, but can stay silent until deformity or growth occurs. 90% of CFD lesions were detected before the age of four [3]. The differential diagnosis of CFD includes benign fibro-osseous lesions of the jaws, giant cell tumors, aneurysmal and simple bone cysts. They have common histologic findings such as hyperproliferation of fibrous material mixed with bony structures and irregular bone [4].

Case presentation

A 11-year-old girl complaining of facial asymmetry and right maxillary prominence was referred to our practice. She had no history of previous trauma or intervention.

MRI examination of the facial bones was requested for this girl. The examination was performed using Toshiba MRI scanner (Toshiba Medical System, New York, USA) and the usual routine sequences were obtained. The hallmark of the examination was partial opacification of the right maxillary sinus by virtue of ill defined intensity eliciting heterogeneous intermediate T1 and predominantly low T2 signals (Figure 1). Complementary CT scan was then performed using Aquillion One CT scanner (Toshiba Medical System, New York, USA), with tube potential set at 110 Kv, current at 320 mA, collimation at 1 mm and table movement at 1 mm/s. An attached workstation and software were utilized to reconstruct axial source images into 2D multiplanar coronal images. The previously thought to be partial opacification of the sinus appeared to be expansion of the right maxillary bone itself with secondary encroachment on the sinus. The right maxillary bone was expanded showing ground glass opacities, however the overlying cortex was completely intact with no evidence of fracture or breaching (Figure 2).

Correspondence to: Tamer W. Kassem, Assistant Professor, Department of Diagnostic and Interventional Radiology, University of Cairo, El-Manial Street, Cairo University Hospitals (Kasr El Ainy), Faculty of Medicine, Egypt, Tel. +201223130682; Fax: 002-02-23634717; Email: tamerwahid@hotmail.com

Key words: fibrous dysplasia, right maxillary

Received: October 10, 2017; Accepted: October 30, 2017; Published: November 02, 2017
The diagnosis of fibrous dysplasia was made based on CT criteria of the lesion and was confirmed lately by histopathological examination. The current case shows to what extent MRI findings could be deceiving in interpretation of bony lesions particularly in facial region and confirm the utmost importance of CT scan in assessment of craniofacial bony diseases.

Conflicts of interest

The authors declare no conflict of interests.

References