Dental and Facial Bone Abnormalities in Pyknodysostosis: CT Findings

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SUMMARY: Pyknodysostosis is an autosomal-recessive disorder of osteoclast dysfunction causing osteosclerosis, with associated maxillofacial anomalies. Multidetector CT with multiplanar and 3D reconstructions illustrated the pathologic findings in this case. Abnormalities included multiple retained deciduous teeth, unerupted teeth with associated follicles, an irregularly expanded alveolus and body of the mandible, and an obtuse mandibular angle. Volume-rendered imaging better delineated the irregular dentition, with crowding and retention of deciduous teeth.

CASE REPORT

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Pyknodysostosis is a rare autosomal-recessive disorder of osteoclast dysfunction causing osteosclerosis. The name derives from the Greek “pyknos,” meaning dense. The disorder is also known as Toulouse-Lautrec syndrome, named for the famous French artist who was thought to be afflicted with pyknodysostosis.

Case Report

A 20-year-old man with pyknodysostosis reported for a CT of the face, requested for preoperative planning. The patient had presented to the oral-maxillofacial surgery clinic with the complaint, “I need some teeth.” Panorex radiograph revealed multiple disorganized crowded deciduous and permanent teeth within the expanded maxillary and mandibular alveolar processes (Fig 1A). Many of the teeth were unerupted or only partially erupted. Precise evaluation of anatomic relationships among teeth was difficult secondary to overlap. This problem was worst in the areas of tooth crowding. Lucencies around the unerupted teeth, possibly representing normal follicles, were present; however, ill-defined borders may have indicated a superimposed inflammatory/infectious process.

Noncontrast CT of the maxillofacial bones was performed on a 4-detector-row CT scanner (Phillips MX-8000, Phillips Medical Systems, Andover, Mass). Axial images of the face were obtained by using 1.3-mm collimation, and multiplanar and 3D reconstructions were generated on a workstation (Voxar, Framingham, Mass). The scout topogram revealed Wormian bones within the lambdoidal suture, a finding often associated with pyknodysostosis (Fig 1B). Axial (Fig 1C, -D), coronal (Fig 1E), and sagittal (Fig 1F) reformatted images demonstrated multiple dental abnormalities. There was persistence of the deciduous teeth within the maxilla and mandible, causing marked crowding. The teeth were misaligned and disorganized, and multiple teeth were unerupted or only minimally erupted. Unerupted teeth were surrounded by well-defined lucent areas, consistent with normal follicles. Poorly marginated lucencies in this location suggested inflammation and/or infection. Along with the dental abnormalities, marked hypoplasia of the maxillary, frontal, and sphenoid sinuses was noted. Unerupted maxillary teeth were seen in the floor of the hypoplastic maxillary sinus, with a few teeth protruding into the antrum (Fig 1E, -F). The hard palate was relatively deep and grooved. The gonial angle of the mandible was obtuse. Note was also made of mild proptosis. Sagittal reformatted images revealed a relatively beaked nose. The surface-rendered 3D image (Fig 1G) gave an overview of these deformities. Irregular dentition was best delineated with volume-rendered display (Fig 1H). Interactive review at the workstation with rotation in any axis and simultaneous orthogonal/oblique multiplanar reconstructions allowed a thorough understanding of all the abnormalities.

Discussion

Pyknodysostosis is an autosomal-recessive disorder of osteoclast dysfunction causing osteosclerosis. Some features of pyknodysostosis overlap the more common osteopetrosis and cleidocranial dysostosis. It is believed that the first case description was in 1923 by Montanari; however, it was not until 1962 that Maroteaux and Lamy defined the characteristic features of pyknodysostosis. General features include short stature (<150 cm), generalized diffuse osteosclerosis with a tendency for fracture after minimal trauma, hypoplastic clavicles, as well as acro-osteolysis with sclerosis of the terminal phalanges—a feature that is considered essentially pathognomonic. Cranial and maxillofacial features include frontoparietal bossing, thick calvaria, open fontanelles and sutures, hypoplastic paranasal sinuses, Wormian bones in the lambdoidal region, relative proptosis, beaked nose, and an obtuse mandibular gonial angle, often with relative prognathism. These findings were seen in our patient, with hypoplastic sinuses being a prominent feature.

Patients may present with frequent fractures, recurrent dental abscesses, or obstructive sleep apnea. Parental consanguinity is recognized as a cause of this autosomal-recessive disorder, the responsible gene being located on chromosome 1q21. This gene encodes cathepsin K, a cysteine protease that is expressed in normal osteoclasts and is mutated in patients with pyknodysostosis. Intracranial features include persistence of deciduous teeth, with premature or delayed rupture of permanent teeth, which can cause crowding. Also seen are tooth misalignment, enamel hypoplasia, and a grooved palate. Dental abnormality was the most impressive finding in this patient, with persistence of deciduous teeth and delayed rupture of permanent teeth. As mentioned previously, the follicles of these unerupted teeth may become infected, leading to abscess formation.

There is limited published material describing the CT findings in patients with pyknodysostosis. CT findings including hypoplastic sinuses, poor dentition, and thickening of the calvaria have been described for osteopetrosis—a similar more common entity. Compared with Panorex radiographs, CT with reconstructions demonstrates greater anatomic detail. Axial imaging defines exact relationships between the unerupted teeth not possible on Panorex alone. Coronal and sagittal reformattting facilitates interpretation of anatomic relationships in the craniocaudal direction. Surface rendering gives an overall perspective of the underlying deformities, while multiplanar imaging and reconstructions facilitated evaluation of the underlying bone abnormalities.
bony abnormalities, and volume rendering aids in determining the 3D relationships among the abnormal teeth by making the alveolar process less conspicuous. Demonstrating the exact relationships among abnormal teeth can aid the surgeon in planning extractions and/or reconstructions. Also, improved visualization of cortical bone can aid in distinguishing infected follicles and dental abscesses from the normal well-corticated follicles of unerupted teeth. This helps the surgeon gain a more complete un-
standing of the abnormal anatomy and is useful for planning of tooth extraction and/or implantation of prosthetics. MR imaging findings in patients with pyknodysostosis generally reveal normal cortical thickness in the calvaria; however, there is increased trabecular bone within the medullary cavity, which causes decreased marrow signal intensity.9

In summary, axial multidector CT with multiplanar reformatted 3D surface-shaded and volume-rendered images demonstrated detailed anatomic and pathologic features of the teeth and facial bones in this patient with pyknodysostosis. Excellent delineation of unerupted and partially erupted teeth was obtained.

References