Non syndromic generalised macrodontia

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Abstract

The purpose of this report is to present a case of non-syndromic generalized macrodontia. Macrodontia is the term used to describe teeth which are larger than normal. Three types of macrodontia are recognized: true generalized macrodontia, relative generalized macrodontia and macrodontia involving a single tooth. Literature review reveals that generalized macrodontia is usually associated with syndromes. Macrodontia may be associated with other syndromes; early recognition of this anomaly is essential to provide proper management. There are many cases reported with macrodontia associated with syndrome, however, our case is particularly unique as it had generalized macrodontia, and was not associated with any syndrome. Macrodontia may cause problems with aesthetics and also crowding if there is discrepancy between the dimensions of the teeth and size of the jaw treatment.

Key words: Macrodontia, True generalized, Associated with syndromes

Introduction

Variation in size, form and morphology of the teeth result in anomalies such as macrodontia, microdontia, hyperdontia, hypodontia, taurodontism and dens in dente. Macrodontia or megadontia refers to teeth that appear larger than the normal size. Some or all teeth may be affected. Macrodontia can be classified as three types: true generalized macrodontia, relative generalized macrodontia and macrodontia involving a single tooth1.

i) True generalized macrodontia - several teeth are larger than normal. Seen in pituitary gigantism and pineal hyperplasia with gigantism.

ii) Relative generalized macrodontia - teeth are normal or slightly larger than normal in small jaws.

iii) Macrodontia of single teeth - this is relatively uncommon. The tooth may appear normal in every aspect except for its size. True macrodontia of a single tooth should not be confused with fusion of teeth, in which, the union of 2 or more teeth results in a single large tooth, early in odontogenesis2.

Case report

A 23 yr old female patient (Fig 1) visited Department of Oral Medicine and Radiology with a chief complaint of a decayed lower left back tooth since 1 year. The tooth was asymptomatic. Her medical history and family history were unremarkable. Our patient was a full term baby and no genetic disorders were found in the family history. Results of physical examination and routine laboratory tests were within normal limits. She was moderately built and well nourished. Her vital signs were stable.

On extra-oral examination, no abnormalities were detected. General examination revealed no associated features like facial dysmorphism, abnormal hair implantation, Cutaneous syndactyly, webbed short neck, cryptorchidism and mental retardation.

On intra-oral examination, gingival bleeding was appreciated on probing. Dental caries was present with respect to 17,11,21,26,36,37,45,46,48 and 13 was buccally placed. Generalized macrodontia (Fig 2) was present along with Talon’s cusp with respect to 11 and 21 (Fig 3). A working diagnosis of True Generalized macrodontia was given. However various syndromes are associated with generalized macrodontia, the features of which are summarized in Table 1.
Full mouth intra oral periapical radiograph was taken which showed teeth larger than normal (Fig 4). Radiographic measurement of the mesiodistal dimension of the various teeth are depicted in Table 2.

As of date, the term “large” is still controversial and subjective. Generalized macrodontia is rare and tooth width also varies between males and females. In our case, the width of the central incisors was 12mm, however literature reveals that the average width of central incisors is 8-8.5mm. The patient's other teeth were also larger than normal.

Discussion

Macrodontia of single tooth is relatively uncommon. Isolated teeth displaying macrodontia can result from twinning abnormalities that originate during the proliferation phase of development. Hemangioma/Lymphangioma can result in an increase in the size and advanced development of adjacent teeth. Fusion and gemination are the most common twinning abnormalities, and both demonstrate enlarged crowns. Males demonstrate a higher frequency of macrodontia and hyperdontia, while females have a greater prevalence of microdontia and hypodontia. The aetiology of macrodontia is unknown, but genetic and environmental causes have been proposed. Generalized macrodontia has been associated with chromosomal anomalies like XYY males (Klinefelter syndrome) and endocrine disorders like pituitary gigantism and insulin-resistant diabetes. Localized macrodontia has been associated with syndromes like the otodental syndrome especially in the posterior segment and patients with unilateral facial hyperplasia who exhibit macrodontia on the affected side. In our case we found enlarged teeth in normal-size jaws, so it can be classified as true generalized macrodontia which was not associated with any syndrome. Radiographic features include an increased size in both erupted and unerupted teeth, the shape of the tooth is usually normal, but some teeth may show mildly distorted morphology.

Generalized macrodontia may pose serious aesthetics and functional disturbances such as malocclusion, crowding and disfigurement. Orthodontic treatment can be accomplished if arch length discrepancy develops. Crown and bridge prosthodontic reconstruction may help to achieve an esthetic appearance. Orthodontic correction of the malocclusion was planned in our patient.

Though we report a case of True Generalised Macrodontia without syndromic association, practitioners should be aware of various syndromes that exhibit True...
Generalised Macrodontia. Recognizing such syndromes early will help in planning appropriate dental and medical care to such patients.

Table 1: Syndromes Associated With Macrodontia

<table>
<thead>
<tr>
<th>Syndromes</th>
<th>Clinical features</th>
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<tr>
<td>Otodental syndrome</td>
<td>Autosomal dominant. Abnormalities of deciduous and permanent dentition included bulbous canines, globe-shaped posterior teeth and agenesis of maxillary premolars. High-frequency sensorineural hearing loss, generalized macrodontia, delayed mineralization of the mandibular premolars, and supplementary permanent maxillary canines.</td>
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<td>Schinzel-Giedion syndrome</td>
<td>The orofacial features include coarse facies, frontal bossing, ocular hypertelorism, anterior open bite and macrodontia.</td>
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<td>Dubowitz syndrome</td>
<td>Rare condition. Affected individual presents with dysmorphic facial features and manifests growth retardation. Dental features include macrodontia, hypodontia, delayed eruption, and midline diastema.</td>
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<tr>
<td>Cockayne’s syndrome</td>
<td>Patients with this syndrome present failure to thrive, short stature, premature aging, neurological alterations, photosensitivity, delayed eruption of the primary teeth, congenitally absent of some permanent teeth, partial macrodontia, atrophy of the alveolar process and caries.</td>
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<tr>
<td>KBG syndrome</td>
<td>Rare condition characterised by a typical facial dysmorphism, macrodontia of the upper central incisors, skeletal (mainly costovertebral) anomalies and developmental delay, short stature, electroencephalogram (EEG) anomalies (with or without seizures) and abnormal hair implantation. Cutaneous syndactyly, webbed short neck, cryptorchidism, hearing loss, palatal defects, strabismus and congenital heart defects are less common findings.</td>
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Table 2: Radiographic measurement of the mesiodistal dimension of various teeth

<table>
<thead>
<tr>
<th>Right and left</th>
<th>In the present Case</th>
<th>Average width</th>
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<tbody>
<tr>
<td>Maxillary Central Incisor</td>
<td>11.1mm</td>
<td>8 mm</td>
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<tr>
<td>Maxillary Premolar</td>
<td>9mm</td>
<td>7mm</td>
</tr>
<tr>
<td>Mandibular Central Incisor</td>
<td>6mm</td>
<td>5mm</td>
</tr>
<tr>
<td>Mandibular Lateral Incisor</td>
<td>6mm</td>
<td>4mm</td>
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<tr>
<td>Mandibular first molar</td>
<td>12 mm</td>
<td>10mm</td>
</tr>
</tbody>
</table>

References


