Orofaciodigital syndrome with mesomelic limb shortening

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SUMMARY Two sisters, the children of first cousin Pakistani Moslem parents, have unusual facies, tongue hamartomata, pre- and postaxial polydactyly, severe talipes, and mesomelic limb shortening associated with tibial dysplasia. Homozygosity for a recessive gene defect is probable. The phenotype resembles, but is distinct from, the orofaciodigital syndromes delineated to date. We suggest that this condition be labelled OFD IV.

Orofaciodigital (OFD) syndrome is characterised by a median pseudo-cleft of the lip, cleft palate, tongue lobulation and hamartomata, multiple oral frenulae, a flat face, hypoplastic zygomatic arch, and both pre- and postaxial polydactyly. Milia of the face, the absence of deafness, and bilateral preaxial polydactyly are features which help to distinguish type I from type II (Mohr), but the principal basis for this distinction is the differing patterns of inheritance. Type I behaves as an X linked dominant, lethal in the hemizygous male, while type II is autosomal recessive. In 1971 Sugarman et al. reported a sib pair with 'jaw winking', supernumerary teeth, polydactyly, and mental retardation. This syndrome has been designated OFD III.

Abnormalities of the long bones are not a recognised feature of the OFD syndromes. Rimoin and Edgerton in 1967 described two sibs with features of OFD II and abnormal proximal metaphyses of the tibiae and fibulae. A third sib, with similar digital features, was stillborn. Goldstein and Medina reported a sib pair with features of type II OFD syndrome, drawing attention to the dental anomalies. Both children were below the third centile for height, with mesomelic shortening in all four limbs. Radiographs revealed flared metaphyses in the long bones of the forearm and foreleg with abnormal constriction of the metaphyses of the proximal tibiae. Temtamy and McKusick, noting the earlier reports, described a child with the facial features of OFD, whose thorax was small and whose limbs were short, with, in particular, rounded proximal tibial metaphyses. They labelled this child and another with a hypoplastic epiglottis as examples of a Mohr-Majewski compound. The Majewski syndrome is a lethal recessive disorder in the short rib polydactyly category. Severe dysplasia of the tibiae and hypoplasia of the epiglottis are characteristic features.

Using the same eponym, Baraitser et al. recently reported the older of the sibs described here, and suggested that this might represent a distinct recessive entity. We report a second affected child in the same sibship with the same features and review the classification of this syndrome.

Case report

III:2, who was the subject of an earlier report, was born on 9.9.81 to first cousin Pakistani Moslem parents. The parents are first cousins. In 1983 a similar pregnancy miscarried at 7 months. A stillborn was born after the next pregnancy in 1984. The family tree is shown in Figure 1.

FIG 1 Family pedigree.

Received for publication 26 October 1983.
Accepted for publication 29 November 1983.
TABLE  Clinical features of the patients.

<table>
<thead>
<tr>
<th></th>
<th>III.2</th>
<th>III.3</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Gestation</strong></td>
<td>33 weeks</td>
<td>40 weeks</td>
</tr>
<tr>
<td><strong>Birth weight</strong></td>
<td>1520 g</td>
<td>2870 g</td>
</tr>
<tr>
<td><strong>Facial features</strong></td>
<td>Low set ears</td>
<td>Low set ears</td>
</tr>
<tr>
<td></td>
<td>Micrognathia</td>
<td>Micrognathia</td>
</tr>
<tr>
<td><strong>Oral features</strong></td>
<td>Posterior cleft plate</td>
<td>High intact palate</td>
</tr>
<tr>
<td></td>
<td>Sublingual hamartoma</td>
<td>Three sublingual hamartoma</td>
</tr>
<tr>
<td><strong>Ocular features</strong></td>
<td>Mild proptosis</td>
<td>Mild proptosis</td>
</tr>
<tr>
<td></td>
<td>Oculomotor apraxia</td>
<td>Oculomotor apraxia</td>
</tr>
<tr>
<td><strong>Skeletal features</strong></td>
<td>Hands: postaxial polysyndactyly</td>
<td>Hands: postaxial polysyndactyly</td>
</tr>
<tr>
<td></td>
<td>Feet: pre- and postaxial polysyndactyly</td>
<td>Feet: pre- and postaxial polysyndactyly</td>
</tr>
<tr>
<td></td>
<td>Severe talipes equinovarus</td>
<td>Severe talipes equinovarus</td>
</tr>
<tr>
<td></td>
<td>Tibial dysplasia and generalised mesomelia</td>
<td>Tibial dysplasia and generalised mesomelia</td>
</tr>
<tr>
<td><strong>Progress and development</strong></td>
<td>Recurrent chest infections</td>
<td>Recurrent chest infections</td>
</tr>
<tr>
<td></td>
<td>Failure to thrive</td>
<td>Gastro-oesophageal reflux requiring fundoplication</td>
</tr>
<tr>
<td></td>
<td>Delayed milestones</td>
<td>Failure to thrive</td>
</tr>
<tr>
<td></td>
<td>Bilateral congenital conductive hearing loss</td>
<td>Delayed milestones (?related to illness)</td>
</tr>
<tr>
<td></td>
<td>CT scan: mild cerebral atrophy</td>
<td>Bilateral congenital conductive hearing loss</td>
</tr>
</tbody>
</table>

By the time a diagnosis was reached in this child, the mother was pregnant again. The second affected sib was born on 7.9.82. Fig 1 shows the pedigree and the table summarises the clinical features. Fig 2 shows the clinical and radiological features in III.2. Fig 3 shows the clinical and radiological features in the second affected sib III.3.

FIG 2 First affected child III.2.
(a) Facial appearance.
(b) Sublingual hamartomata.
(c) AP x-ray of tibiae, fibulae, and feet. Symmetrical tibial shortening is present with mild lateral bowing of the mid shafts. The fibulae are long in relation to the tibiae and there are dislocations at knees and ankles. The right foot shows polysyndactyly (eight digits). The first metatarsal is short, broad, and rounded and the duplicated phalanges are correspondingly wide. Postaxial polydactyly is also present. The left foot shows similar changes but there is also virtually no ossification of several middle and terminal phalanges.
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(d) PA x-ray of both hands. The right hand shows postaxial polydactyly (seven digits but only five metacarpals). Abnormal pseudoepiphyses are associated with the metacarpals. The index finger shows camptodactyly with shortening of the middle phalanx. The left hand also shows postaxial polydactyly (six digits but five metacarpals). The middle phalanges of the index and little fingers are abnormally short.

(e) Lateral x-ray of knees. Dislocation and proximal shortening of both tibiae.

Oculomotor apraxia is a defect of voluntary and optically induced horizontal eye movement with retention of voluntary vertical gaze. It may be associated with absence of the corpus callosum though this was not evident on the CT scan of III.2.

Discussion

The occurrence of the same pattern of malformation in female sibs is suggestive of autosomal recessive inheritance. When the parents are healthy first cousins this genetic basis becomes very probable, even though the parents come from a group where inbreeding is frequent.

The mesomelic limb shortening, particularly involving the tibiae, is of importance in classification, antenatal diagnosis, and prognosis. The children in this report have many of the features of the OFD syndromes, but the specific defects of the long bones justify the separate designation of an OFD syndrome with mesomelia: OFD IV.*

*These children were presented (by CMH) at the European Society of Paediatric Radiology, Paris 1983. The proposed designation OFD IV was accepted.
We are grateful to Dr M O Savage for permission to publish this case and to Mrs Melanie Barham and Ms Carol Reeves for preparation of the manuscript and illustrations respectively.

References


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doi: 10.1136/jmg.21.3.189

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