Nager acrofacial dysostosis: a rare genetic disorder causing bilateral temperomandibular joint ankylosis in a 10-year-old girl

M Hari Kumar,1,2 M Siva Kumar,3 Vishalakshi Siva Kumar,3 Sabitha Hari Kumar2

1Department of Oral Medicine and Oral Radiology, Rajas Dental College and Hospital, Tirunelveli, Tamil Nadu, India
2Dental Department, Jeba Dental Clinic (a multispecialty Dental Clinic), Nagercoil, Tamil Nadu, India
3Department of Maxillo Facial, Doctor Kamalkanadasan Dental Clinic, Chennai, Tamil Nadu, India

Correspondence to
Dr M Hari Kumar, drhari.omrd@gmail.com

Accepted 5 December 2015

DESCRIPTION

A 10-year-old girl presented to the oral medicine department, with restricted mouth opening since childhood. Her family history revealed that she was the third child born of a consanguineous marriage. She was born after an unremarkable pregnancy, with a birth weight of 2.3 kg. Physical examination revealed downslanting palpebral fissures, microretrognathia, malar hypoplasia, a sloping forehead, high nasal bridge with an upturned nasal tip and short philtrum (figure 1). She was also noted to have short forearms, proximal radioulnar synostosis (figure 2), hypoplastic thumbs and relatively short overlapping toes (figure 3A, B). Oral examination revealed extremely poor oral hygiene with a narrow V-shaped short palate; the mouth opening was 15 mm.

Lateral cephalogram showed an underdeveloped mandible with an extreme posterior inclination of the mandibular base and also complete obliteration of the temperomandibular joint (TMJ) space (figure 4). Panoramic radiographic examination showed reduced vertical height of ramus, and multiple impacted teeth in the maxilla and mandible, with complete obliteration of right and left TMJ space (figure 5). Hearing assessment by brain stem-evoked response audiometry showed a bilateral profound hearing loss. In this case, genetic analysis was not performed due to financial constraint. From the girl’s overall clinical and radiographic features, she was diagnosed as having Nager acrofacial dysostosis. The patient’s mother gave full informed written consent to this report being published.

Nager syndrome was first recognised by Nager and de Reynier in 1948. They used the term acrofacial dysostosis to describe their patient. The syndrome (preaxial acrofacial dysostosis) is due to aberrations in development of the first and second branchial arches and limb buds.1

Nager acrofacial dysostosis is characterised by a variety of craniofacial features including malar hypoplasia, cleft palate and micrognathia. The limb anomalies associated with Nager acrofacial dysostosis are a significant feature of this syndrome and help the clinician to differentiate this entity from other craniofacial syndromes. Patients with Nager syndrome often require several surgical procedures for correction of congenital anomalies, including cleft palate repair and mandibular lengthening with distraction osteogenesis or orthognathic surgery, among others, including bone grafts and ramus reconstruction.2

Figure 1 Downslanting palpebral fissures with microretrognathia and malar hypoplasia.

Clinical management in these cases mainly depends on a multidisciplinary approach focused on treatment of symptoms. Panoramic radiograph and lateral cephalogram obtained during the initial diagnostic procedure revealed bilateral TMJ ankylosis. The patient was referred to the oral and maxillofacial surgery department and was motivated for surgical treatment.

Figure 2 (A and B) Short forearms with proximal radioulnar synostosis.
Total prosthetic TMJ replacement has been successfully used for various pathological, developmental and traumatic conditions, including inflammatory arthritis involving the TMJ, recurrent fibrous or bony ankylosis not responsive to other modalities of treatment.²

In this case, correction of the TMJ ankylosis was planned with a bilateral custom total prosthetic joint replacement; this surgery helps in re-establishing the form and function of the mandible and also significantly improves the patient’s quality of life.

Learning points

▸ Nager acrofacial dysostosis is a rare congenital syndrome and nearly 94 cases have been reported in the medical literature. Nager syndrome is characterised by craniofacial, limb and musculoskeletal anomalies. The orofacial features are similar to those of Treacher Collins syndrome, but it can be distinguished from Treacher Collins syndrome by preaxial limb deformities.

▸ Nager acrofacial dysostosis is associated with various syndromic deformities, making it difficult to predict the treatment result. The clinical management for most cases mainly depends on a multidisciplinary approach consisting of maxillofacial radiology, oral and maxillofacial surgery, paediatrics and neonatology, otolaryngology, anaesthesia, plastic surgery and orthopaedic surgery.

Contributors

MHK and MSK contributed to diagnosis of the patient, concept of the paper, acquisition of the data, and drafting, revision and final approval of the article. VSK contributed to diagnosis of the patient, concept of the paper, acquisition of the data, and revision and final approval of the article. SHK contributed to diagnosis of the patient, concept of the paper, and drafting, revision and final approval of the article.

Competing interests

None declared.

Patient consent

Obtained.

Provenance and peer review

Not commissioned; externally peer reviewed.

REFERENCES

