

# **Cleidocranial Dysplasia - A case report**

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**Vol. 17, No. 2 (2006-05 - 2006-08)**

Biomedical Research 2006; 17 (2): 129-132

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**Key Words:** Cleidocranial Dysplasia, Supernumerary teeth, Clavicle, Fontanel, Coronoid process, Mandibular notch.

Accepted March 06, 2006

## **Abstract**

Cleidocranial Dysplasia is a rare autosomal dominant disorder affecting skull, jaws, teeth, clavicle along with other skeletal abnormalities. It is noteworthy that the case of a 28-year-old male reported here concerns the dental surgeon as it deals with high number of impacted and supernumerary teeth as well as additional unusual findings such as elongation of coronoid process and rounded gonial angle that make it quite peculiar.

## **Introduction**

Cleidocranial Dysplasia also known as Marie and Sainton's disease is a rare autosomal dominant disorder affecting bones and teeth [1,4]. It affects men and women with equal frequency [1,2,4] and is characterized by abnormalities of skull, dentition, jaws and clavicle. It is seen that the fontanel are open despite the age of the patient being far beyond the normal closure time [1,2,3-6]. In our patient there were 64 teeth and such a high number has been reported in rare instances along with deepening of mandibular notch. In addition to this concave profile due to under developed maxilla [9], a brachycephalic skull has also been reported [7,9]. The bridge of the nose was broad and depressed. The mandible appears to be prognathic with a depressed forehead.

## **Case Report**

A 28-year-old Asian male of Indian origin was reported to the Department of Oral Medicine and Radiology of Jaipur Dental college, University of Rajasthan, on June 15, 2005. His chief complaint was missing teeth and he desired removable prosthesis. He gave the history of falling of teeth few years back. Upon general examination, he was of average height and his gait showed deviation from normal. It was found that he had

depression in forehead and it extended as a groove between two parietal bones up to the occiput (Fig. 1). The skull was brachycephalic. The midface was depressed because of underdeveloped maxilla, so the mandible appeared prognathic (Fig. 2). The patient gave history that his mother had same kind of scalp and facial profile. The patient had abnormal movement of right shoulder and the right side of clavicle exhibited mobility on manipulation (Fig.1). His long bones of the limbs & bones of digits showed bowing (Fig. 3).

His intra oral examination revealed presence of the following teeth in the oral

87654 DC	123 DE 67,
654 1	12 CD 567

cavity,

of which root stumps of  $\begin{array}{c|c} 6 & E \\ \hline & C \end{array}$  were seen

The supernumerary teeth seen in between  $\begin{array}{c} | \\ - \\ 23 \end{array}$  region, and  $\begin{array}{c} | \\ + \\ 1 \end{array}$  were peg-shaped (Fig.4).  
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Further the orthopantomogram outlined the presence of total 64 teeth in both the jaws. Out of these, 39 were im-pacted and supernumerary and 25 were seen in the oral cavity. The mandibular notch showed excessive deepening causing the elongated appearance of coronoid process (Fig. 5).

The gonial angle appeared extremely rounded and lack of definition of the neck of condyle (Figs. 5,7).

In lateral skull radiograph, there was depression in anterior fontanel region. Both frontal bones overlapped giving rise to appearance of double density. Posterior fontanel region also open as seen in radiograph. The lambdoid sut-ure region showed wormian bones (Fig. 6,9). The hypoplastic appearance of maxilla was clearly evidenced on the lateral skull radiograph and lateral cephalogram (Figs. 6,7).



Fig. 1: The Patient of Cleidocranial dysplasia Showing hypermobility of right shoulder and depressed forehead.



Fig. 2: Patient's profile showing concavity due to underdeveloped maxilla.



Fig. 3: Patient shows bowing of the bones of hands and feet.



Fig. 4: Photograph of Intra oral view of the patient.



(For larger image, click [here](#))

Fig. 5: Panoramic radiograph shows remarkable number of Supernumerary and malposed teeth, deepening of mandibular notch and lack of definition of neck of condyle.



Fig. 6: In Lateral skull view depression in anterior fontanel and appearance of double density because of overlapped frontal bones, wormian bones also seen



Fig. 7: In Lateral cephalogram, underdeveloped maxilla, Deep mandibular notch and rounded gonial angle seen Cleidocranial Dysplasia – A case report



Fig. 8. Chest radiograph showing hypoplastic appearance of the clavicles, they were not angulated as normally seen and scapulae were laterally placed.



Fig. 9: P.A. skull radiograph shows open anterior fontanel and sutures of skull.

In posteroanterior radiographic view of chest the scapulae were laterally placed, clavicles were not angulated as normally seen. The right side of clavicle appeared hypoplastic as compared to left side (Fig.8).

## **Discussion**

Our observations in this case are the presence of excessive number of supernumerary teeth in the maxillary molar region and mandibular premolar region in Cleidocranial Dysplasia syndrome [2,9,10]. The latter resemble the premolar teeth. The total number of teeth were 64, that was in extremely high number [2,9-11]. As mentioned in previous cases, the fontanels were also open although the patient was in his late twenty years [1,5,7,8] i.e. 28 years old.

Along with this, an unusual finding encountered was the deepening of mandibular notch hence elongated appearance of coronoid process of the mandible. The gonial angle was rounded off. There was lack of definition of the neck of condyle. The normal constricted appearance of condyle was absent. In this case the clavicles were present but not formed to the full extent, the hypermobility was found in the right shoulder to a greater extent as compared to the left. [8,10].

The presence of unerupted teeth interfered with day to day dental care of the adult patient. It is often required to fabricate prosthesis over unerupted teeth [12].

## **Conclusion**

This article presents a 28 year old male suffering from Cleidocranial Dysplasia. This patient refused an elaborate surgical correction. In this subject fabrication of prosthesis was carried out to replace the missing dentition and advised about the importance of oral preventive care in daily routine. Such patients should be diagnosed at the early mixed dentition stage to render treatment at suitable interval in the form of oro-surgical and orthodontic correction, so that such problems are overcome before adulthood is attained. The followup is essential in such a syndrome in order to check the development of odontogenic cysts etc.

This peculiar case has been reported to focus on the unusual findings so that the medical and dental professionals become familiarize with such features on confronting subjects with Cleidocranial Dysplasia.

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