Untreated craniostenosis causing severe mental retardation with X-ray skull showing silver beaten appearance skull on: Role of physician in early diagnosis and surgical management

Guru Dutta Satyarthee* and Satyajit Panda

1Associate professor, Department of Neurosurgery, Gamma Knife, AIIMS, New Delhi, India
2Senior resident, Department of Neurosurgery, Gamma Knife, AIIMS, New Delhi, India

Abstract

Craniosynostosis is rare but important clinical condition characterized by early fusion of skull suture causing restriction of growth of developing brain and ultimately arrest of brain growth. It is a cause of acquired mental retardation, which can be avoided and prevented by early diagnosis and early surgical intervention as delay causes irreversible loss of vision and permanent cognitive impairment. So all physicians should be made aware of this rare congenital deformity. Authors report a neglected case of craniostenosis, presented with deformity of head with mental retardation, which was operated after delay lead to poor neurological outcome.

Introduction

Craniosynostosis is characterized by early fission of sutures of skull, when premature closure of the sagittal suture occurs called as Scaphocephaly [1]. It causes arrest of growth of brain and causes deformity of skull along with features of raised intracranial pressure. However, early surgical intervention can lead to normal growth [2]. However, neglected cases can develop typical scaphoid appearance of head and silver beaten appearance or also known as beaten brass skull, refers to the prominence of gyral impressions on the inner table of the skull observed throughout the skull vault and typically associated with raised intracranial pressure in children [3-5]. However, convolutional markings may be normal, however, such convolutions are usually confined to the posterior part of the inner table of the skull. Every case of craniostenosis should be subjected to surgical intervention at as early as possible preferably within first three months of infancy to protect from bad effect of raised Intracranial pressure [6-8].

Case illustration

A 7-year-old male boy presented to our outpatient clinic with complaints of abnormal shape of head along with abnormal facial profile. He was mentally subnormal and also had abnormal protrusion of both eyes associated with bilateral progressive diminution of vision. On examination, patient had bilateral proptosis. Visual acuity was finger counting at 3 feet in both eyes. There was a palpable ridge over sagittal suture line suggestive of fused suture. He had head circumference of 47 cm with biparietal diameter of 17 cm. Non-contrast computerised tomography head with three-dimensional reconstruction revealed a small neurocranium with fusion of all calvarial sutures (Figure 1).

Digital X-ray skull showed heavy convolutional markings as a result of long-standing raised intracranial pressure giving a classical “Silver-beaten appearance of skull” (Figure 2,3). Patient was taken up for surgery. Strip cranietomy with barrel-stave osteotomy was done along with lax duraplasty. Patient had uneventful post-operative course and was doing well at follow-up visit six months after surgery.

Discussion

Scaphocephaly is a type of craniosynostosis, characterized by premature fusion of sagittal suture and causes deformed appearance of skull resembling keel of a boat as long and thin skull. It represents most common type of craniosynostosis, constituting more than half of all cases [2]. The Scaphocephaly is usually isolated and usually not associated with syndromes, but observed in premature infants.

As growth of the skull bones is prevented by fused sagittal suture, however compensated by expansion at lambdoid the coronal sutures [3]. The skull appears as elongated and shortened in the biparietal diameter and palpable ridge of the sagittal suture as compensated frontal and occipital bones for the restricted lateral expansion of the parietal bones. It is associated with closed fontanelle [5].

X-ray skull shows presence of elongated skull. It may also show presence of “silver-beaten cranium” is also called as “beaten copper cranium”, which is postulated to be due to indentations in the bone from the underlying gyri due raised intracranial pressure and cerebrospinal fluid pulsations [1]. The unyielding nature of calvarium in craniosynostosis produces a characteristic mottled appearance of

Correspondence to: Guru Dutta Satyarthee, Associate Professor, Department of Neurosurgery, Room no. 714, Neurosciences Centre, AIIMS New Delhi, India, E-mail: duttaguru2002@yahoo.com

Key words: Craniosynostosis, silver-beaten appearance, mental retardation, craniolacunae, management

Received: January 14, 2017; Accepted: February 16, 2017; Published: February 20, 2017
Untreated craniosynostosis causing severe mental retardation with X-ray skull showing silver beaten appearance skull on: Role of physician in early diagnosis and surgical management

Satyarthee GD (2017) Untreated craniosynostosis causing severe mental retardation with X-ray skull showing silver beaten appearance skull on: Role of physician in early diagnosis and surgical management

Gen Int Med Clin Innov, 2017

doi: 10.15761/GIMCI.1000135

The bone with lucencies of varying depth having round and poorly marginated edges.

The pathological mechanism of silver-beaten cranium is caused by sustained high intracranial pressure is causing increased inner table convolutional markings primary cause of copper beaten skull includes craniosynostosis, obstructive congenital hydrocephalus, hypophosphatasia and intracranial tumours.

The silver beaten appearance on imaging study needs different ion from convolution marking and lacunar skull [4,5]. The lacunar skull is also known as craniolacunae represents dysplasia of the membranous skull vault. Lacunar skull is associated with 70% cases Chiari malformation and typically inner table is more affected than the outer with intervening areas of apparent thinning. Lacunar skull appearance represents abnormal collagen development and ossification of membranous bone [2,4,5]. Typical radiological feature of lacunar skull is presence of a groups of oval or round pits on the inner surface of the vault, separated by ridges of bone and usually have predilection for thickest part of frontal, parietal and upper occipital bone [4,6].

The cranial computed tomography confirms the diagnosis of scaphocephaly. It can also show associated microcephaly and other associated pathology, like hydrocephalus, or chinked ventricle with effacement of arachnoid spaces [7,8].

Treatment of craniosynostosis is surgery [2]. The surgical intervention should be carried out the earliest possible occasion as progressive raised intracranial pressure and cognitive impairment can be avoided at early surgery with good neurological outcome and leads to significant improvement in the head shape and allows permit re-growth and development of brain [3].

Conclusion

Early surgery can spare from effect of prolonged raised intracranial pressure and help in scholastic improvement. However neglected cases can develop typical scaphoid appearance of head and silver beaten appearance. Every case of craniosynostosis should undergo surgery at infancy stage itself.

References

1. Tuite GF, Evanson J, Chong WK, Thompson DN, Harkness WF, et al. (1996) The beaten copper cranium: a correlation between intracranial pressure, cranial radiographs, and computed tomographic scans in children with craniosynostosis. Neurosurgery 39: 691-699.
2. Ciurea AV, Toader C, Mihalache C (2011) Actual concepts in scaphocephaly: (an experience of 98 cases). J Med Life 4: 424-431. [Crossref]
3. Farhad Afsah. The Craniosynostoses: Causes, Natural History and Management J R Soc Med. 1983 April; 76(4): 335
4. Desai V, Priyadarshini SR, Sharma R (2014) Copper Beaten Skull! Can It be a Usual Appearance? Int J Clin Pediatr Dent. 7: 47-49. [Crossref]
5. Rühi FJ1, Nickisch N, Alt KW (2007) A historical case of beaten-copper cranium. J Neumnsurg 106: 71-73. [Crossref]
6. Shahdadpuri R, de Vries B, Pfündt R, de Leeuw N, Reardon W (2008) Pseudoarthrosis of the clavicle and copper beaten skull associated with chromosome 10p11.21p12.1 microdeletion. Am J Med Genet A 146A: 233-237. [Crossref]
7. Agrawal D, Singh PK, Sinha S, Gupta DK, Satyarthee GD, et al. (2015) Remaining unconscious: The burden of traumatic brain injuries in India. J Neurol Sci Rural Pract. 6: 520-522.
8. Satyarthee GD, Mahapatra AK (2015) Giant pediatric glioblastoma multiforme causing primary calvarial erosion and suture diastasis presenting with enlarged head. J Pediatri Neurosci. 10: 290-293

Copyright: ©2017 Satyarthee GD. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.