THE ANTLEY-BIXLER SYNDROME: A CASE REPORT AND RESULT OF RADIOHUMERAL SYNOSTOSIS RESECTION

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ABSTRACT
Antley-Bixler syndrome (ABS) is a rare congenital disorder with multiple skeletal and cartilaginous anomalies that demand orthopedic management. In this report we address the diagnostic features and associated malformations of this syndrome and describe the functional outcome of resection of radio humeral synostosis in a 9 month old girl with Antley-Bixler Syndrome.

KEYWORDS: Antley-Bixler syndrome, Radiohomeral synostosis, Craniosynostosis.

INTRODUCTION
Antley-Bixler Syndrome (ABS) is a rare genetic disorder characterized by distinctive craniofacial and skeletal abnormalities.1,2 It was first reported in 1974 by lachertz et al. and again in 1975 by antley and Bixler. However it was in 1977 that the term Antley-Bixler syndrome was designated to describe the constellation of findings. Over the years what is known as Antley-Bixler syndrome has also been published under the terms such as multisynostotic osteodysgenesis, acrocephalo-synankie, and trapezoidocephaly multiple synostosis syndromes.3

Primary features of Antley-Bixler syndrome includes craniosynostosis, midface hypoplasia with or without choanal stenosis or atresia, radio humeral synostosis, femoral bowing and joint contractures.2,4 Cardiac, urogenital and gastrointestinal anomalies have also been reported including septal defect, hydronephrosis, and fused labial minored and anal arter-sian.3,5 Genital anomalies including fused labia minored, reported as well.2,6

Hosalker Review of literature reveals that 41 cases of ABS have been reported so far.5 Here we report a female patient with skeletal features of ABS and sort term result of radio humeral synostosis resection in that patient.

Case History: A nine month old girl, born to a 23- year-old primigravida mother, presented with flexion contracture of right elbow. Her general health was good. The parents were unrelated and family history was not contributory. The pregnancy was uneventful and the baby was a product of normal vaginal
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delivery at 38th week of gestation. On physical examination, there was evidence of mild brachycephaly (cephalic index: 82), mild midface hypoplasia, frontal bossing and depressed nasal bridge, (Fig-1,2).

Examination of limbs revealed fixed 90 deg flexion contracture of the right elbow and absence of the thumb. Radiographs confirmed right radio humeral synostosis and skeletal abnormality of right hand including absence of 1st metacarp with hypoplasia and fusion of the 4 th-5th metacarp. (Fig-2,3).

Skull X-Ray and computed tomographic scans confirmed synostosis of both coronal sutures. Sonography of abdominal pelvic and echocardiography was unremarkable. Following a routine pediatrician and preanaesthetic evaluation, she was brought to the operating room. Under general anesthesia with lateral approach, resection of radio humeral synostosis and interposition of local soft tissue was performed. Full range of motion of elbow was achieved (Fig-3,4). A posterior splint was applied and physiotherapy was initiated. The range of elbow and forearm motion was normal in follow up visit at 18 months after surgery.

DISCUSSION

The etiopathogenesis of ABS is not clearly recognized. This Syndrome is suggested by most
authors to be inherited as an autosomal recessive trait; however it may also result from sporadic mutations especially in fibroblast growth factor 2 locus.5,7 Mutations in the genes for Fibroblast growth Factor receptors (FGFRs) have been reported to be involved in some craniosynostosis syndromes such as Pfeiffer, Apert, cruzon and Jackson-Weiss Syndromes.8 Fibroblast growth factor 8 (FGF8) protein is probably a ligand for FGFR2, is androgenic, and is expressed in the developing central nervous system, face, limb, Kidneys and gonads.4,9 It plays a key role in both midbrain and limb development and as both FGF8 and FGFR2 are located in chromosome 10q25q26, they may be involved in craniosynostosis.10

Prognosis is guarded in infancy and childhood (59% death rate in the first decade of life), but improves with age. Respiratory distress secondary to choanal atresia or stenosis can be a serious factor for mortality.2,5

Synostosis frequently results in early orthopedic consult as fixed contracture of the affected joints and the surgeons should be aware of the frequent associations with this syndrome. Other orthopedic manifestations of Antley-Bixler Syndrome include femoral bowing, ulnar bowing, and camptodactyly, synostosis of tarsal and carpal bones, clubfoot, vertebral body anomalies and prenatal fracture.5 Cohen pointed out that the combination of craniosynostosis and elbow ankylosis as the cardinal features in this syndrome.1

Our patient had milder features of skull and facial anomalies in comparison to similar reported patients. Also she had brachydactyly and thumb agenesis that to our knowledge has not been reported with this syndrome.

Our report describes an uneventful management of such a patient with respect to anesthesia and outcome of radio humeral synostosis resection. Although resection of radio humeral synostosis has not been encouraged in some reports in literature, we succeeded to achieve satisfactory range of motion of the elbow in our 18 month follow up of this patient after resection of the synostosis.

CONCLUSION

Varying bone and joint anomalies are seen with ABS that need orthopedic assessment. Our patient was without urogenital or cardiac disorder. The resection of radio humeral synostosis had a good result at least in a short term follow up.

REFERENCES