

An Omphalocele, Epispedias, Cleft Palate, Cranial deformity and Facial Dysmorphism: A Case with Midline and Laterality Defects

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ABSTRACT

A 7 year old male child with cleft soft palate, omphalocele, epispedias, posterior prominence of the skull, prominent forehead with high anterior hair line, dextraposition of the heart, right sided inguinal hernia, mental retardation, generalized hypotonia and flexion deformity of both toes and fingers presented to the paediatric clinic, Teaching Hospital Karapitiya, for the follow up management. Furthermore, the child had subtle dysmorphic features including, broad nasal bridge, hypertelorism and low set ears. He was the second child of the family and there were no other family history of congenital anomalies. The karyotype was 46XY. Mutations in chromosome bands 3p12-21, *ZIC3* gene in human X chromosome and Wolf-Hirschhorn syndrome involving heterozygous deletion of 4p16.3 region (4p syndrome) can be presented with above clinical features and it is necessary to investigate the patient further for the genetic involvement.

Key words: omphalocele, epispedias, cleft palate, cranial deformity, facial dysmorphism, *ZIC3* gene, 4p syndrome

INTRODUCTION

Disturbances of the normal asymmetric position of organs or situs inversus, have been described as laterality defects. Congenital anomalies of midline structures such as, esophageal defects, anal defects, neural tube defects, cleft lip and palate are considered as midline defects. Animal model studies and human birth defect registries suggest that the midline and laterality defects are etiologically related (1).

All vertebrates have a body configuration with a midline and left and right halves that are symmetric for most external features. Many internal organs have asymmetric placement in the body with respect to the midline. The heart is positioned in the left side of the thorax with its apex pointed towards the left. The spleen and stomach are placed on the left

side, liver in the right side of the abdomen. Alterations of the left/right (LR) asymmetry may cause alteration of the position of organs (heterotaxia or situs ambiguous) or they may produce reversal of LR organ position (situs inversus). Disturbances of LR asymmetry may also be limited to a single asymmetric organ such as the heart in individuals with isolated dextrocardia or dextraposition of the heart. Collectively, any abnormalities of LR asymmetry are referred to as a laterality defect (1). It has been recognized that there is a genetic basis for some disturbances of LR asymmetry in humans. Although most cases are considered sporadic, families with laterality defects have been shown to have an autosomal dominant, autosomal recessive and X-linked recessive inheritance patterns. Thus,

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multiple genes exist that are responsible for human laterality defects, although mutations of single genes may cause laterality defects that can be seen as Mendelian patterns (2,3). Genes corresponding to the chromosome bands 3p12-3p21 has been described in the literature causing midline and laterality defects. Furthermore, mutations in *ZIC3* gene in human X chromosome have been identified in causing midline and laterality defects (4).

There is clear evidence that model animals with disturbances of LR asymmetry to have midline defects more commonly than expected by chance. Some of these defects include neural tube defects, oral clefts, omphalocele, esophageal atresia/tracheoesophageal fistula, imperforate anus, epispedias conotruncal heart defects, diaphragmatic hernia (5).

Four families with a distinctive malformation syndrome which includes diaphragmatic defects, limb deficiencies and ossification defects of the Skull were also reported (6).

Wolf-Hirschhorn syndrome (WHS) is characterized by typical craniofacial features in infancy consisting of wide bridge of the nose continuing to the forehead, microcephaly, high anterior hairline with prominent glabella, widely spaced eyes, epicanthic folds, high arched eyebrows, short philtrum, downturned corners of the mouth, micrognathia, and poorly formed ears with pits and tags. The diagnosis of Wolf-Hirschhorn syndrome is established by detection of a heterozygous deletion of the Wolf-Hirschhorn syndrome critical region (*WHSCR*) within 4p16.3 at 1.4-1.9 Mb from the terminus (7).

CASE REPORT

Index case was a 7 year old male child with cleft soft palate, Omphalocele, epispedias, right sided inguinal hernia, posterior prominence of the skull, prominent forehead with high anterior hair line, mental retardation, hypotonia, dextraposition of the heart, flexion deformity of both toes and fingers. Furthermore, he has having subtle dysmorphic features such as, broad nasal bridge, hypertelorism and low set ears. He was born at term with a birth weight of 2.4kg. He was presented with global developmental delay. No other respiratory or gastro intestinal defects. He was the second child of the family with no other family history of congenital anomalies. His elder brother was apparently normal. Echo cardiogram of the child revealed dextraposition of the heart and no other congenital cardiac anomalies were present. Ultra sound scan of the abdomen revealed normal. His karyotype was 46XY.

The following differential diagnoses were made:

- 1) Mutations in chromosome bands 3p12-21 causing midline and laterality defects.
- 2) Mutations in *ZIC3* gene in human X chromosome causing midline and laterality defects.
- 3) Wolf-Hirschhorn syndrome involving heterozygous deletion of 4p16.3 region (4p syndrome).

The omphalocele, cleft palate and epispedias were treated by the Pediatric and the Maxio-Facial surgeons at the Teaching Hospital Karapitiya. The child was followed up in the Rheumatology clinic and the Physiotherapy department for his rehabilitation.

DISCUSSION AND CONCLUSION

According to the reported literature there is clear evidence that midline and laterality defects occur together more commonly than expected by chance (5). The midline defects include neural tube defects, oral clefts, omphalocele, esophageal atresia/tracheoesophageal fistula, imperforate anus, epispedias, conotruncal heart defects, diaphragmatic hernia whereas the laterality defects include dextraposition of heart, malrotation of gut and other dextraposition of viscera such as right sided spleen (5). It was stated that the mutations in chromosome bands 3p12-21 causing midline and laterality defects associate together more frequently. Furthermore, it was reported that mutations in *ZIC3* gene in human X chromosome causing midline and laterality defects in some families (1).

The 4p syndrome or Wolf-Hirschhorn syndrome is characterized by typical craniofacial features and other abnormalities in infancy (7).

The index case also has shown some clinical features such as, dextraposition of the heart, omphalocele, epispedias, congenital inguinal hernia, cleft palate, facial dysmorphism and cranial anomalies, in favor of the above mentioned differential diagnosis. Furthermore, his karyotyping was 46 XY.

Molecular genetics investigations are important in this case for diagnosing the condition and to identify the etiology of the anomalies.

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