Omphalocele with Dextrocardia - A Rare Association

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INTRODUCTION

Omphalocele and gastroschisis are the two most common congenital malformations of the anterior abdominal wall. Omphalocele is a midline abdominal wall defect with protrusion of abdominal viscera, covered by a membranous sac, into the base of the umbilical cord. Omphalocele is frequently associated with many other congenital malformations, like cardiac anomalies, ranging at the rate of 15 to 54%. In cardiac anomalies, association of omphalocele with dextrocardia has been noticed in very few literatures only. We describe here such a rare association, with a brief review of the literature.

CASE REPORT

A two day old male neonate presented to our surgical unit with an abdominal wall defect (Figure 1). On examination, the child was morphologically normal except for the omphalocele, his vitals were stable, and on examination of the chest the heart sounds were predominantly heard on the right side. A radiograph of the chest and abdomen showed the heart shadow in the right side, along with the omphalocele sac (Figure 2). An echocardiography was done, which showed dextrocardia of the heart, without other anomalies (Figure 3). The omphalocele sac was excised after opening the sac and freeing the contents which were cecum, ascending colon and distal ileum. Anatomical closure of the anterior abdominal wall was done. There were no complications in post-operative period. At three months follow-up, the child was doing well.

DISCUSSION

Omphalocele is a sporadic abnormality with a birth prevalence of about 1 in 4000. The sac is composed of peritoneum, amnion and Wharton’s jelly. Visceral contents in the sac may include loops of intestine, liver, and stomach. Omphalocele has been found to be isolated or associated with various other malformations, like chromosomal abnormalities, non-chromosomal syndromes (including Beckwith–Wiedemann syndrome, Goltz syndrome, Marshall–Smith syndrome, Meckel–Gruber syndrome, Oto-palato-digital type II syndrome, CHARGE syndrome, and fetal valproate syndrome) malformations (like ectopia cordis, body stalk anomaly, etc.).
Malformations of the cardiovascular system include one of the most common congenital malformations in patients with omphalocele. However, dextrocardia with omphalocele has been reported in very few literatures, even including large series. In Stoll et al’s study, dextrocardia has been found to be associated with 5 out of 68 patients. Ventricular septal defect was the commonest, dextrocardia was the second commonest. In another study by Mayer et al, two out of 28 patients had associated dextrocardia, along with anophthalmia, microcephaly, and trisomy-13; both these children did not survive. In many other studies, dextrocardia has not been mentioned. Common heart diseases that have been found with it include ventricular septal defect, atrial septal defect, Tetralogy of Fallot, hypoplastic left ventricle and coarctation of aorta.

Isolated dextrocardia has not been associated with an increased mortality rate. Dextrocardia in combination with other anomalies (multiple or chromosomal anomalies) definitely has been associated with an increased mortality, and the available literatures show that mortality is mainly attributable to the associated anomalies. Hence, some authors have even considered screening for other congenital malformations as mandatory in patients with omphalocele. This case report highlights isolated dextrocardia as a possible association with omphalocele.
REFERENCES


Shakya et al. Omphalocele with Dextrocardia - A Rare Association