Pentalogy of Cantrell (PC) is an uncommon congenital disorder characterized by severe defects in the chest and abdomen, including abdominal visceral prolapsed via umbilical cord (omphalocele), defect in the lower part of the sternum, defect in the front of the diaphragm, defects in the anterior part of the pericardium, and the ectopiacordis. Here, we report a 2-hour-old girl, weighing 3,500 g, who was referred to Shahid Madani Hospital in Khorramabad due to the large omphalocele on her chest with pulsating mass above it. The baby was the first child of a 24-year-old mother who was born with an uncomplicated vaginal delivery. Very rare cases of PC are born as a term new-born.

Case Report

A 2-hour-old girl, weighing 3,500 g, was referred to Shahid Madani Hospital in Khorramabad due to the large omphalocele on her chest with pulsating mass above it. The baby was the first child of a 24-year-old mother who was born with an uncomplicated vaginal delivery (►Video 1).

Video 1

Video showing case report from different positions. Online content including video sequences viewable at: https://www.thieme-connect.com/products/ejournals/html/10.1055/s-0039-1683936.

Her parents were unrelated. Her mothers' pregnancy was uncomplicated and the mother had no health care and medical care during pregnancy. The family history of the disease was negative. The history of abortion, fetal death, and polyhydramnios in the mother was negative. The history of mother's contact with chemicals, toxins, vegetable pests, opium, cocaine, medications (sodium valproate, phenytoin, lithium, lead) radiation, or and environmental factors were negative. In physical examination, the neonatal reflexes were reduced.
The vital signs of the neonate were: blood pressure 45/30, respiratory rate 72 beat/min, temperature 36°C, heart rate 135/min and SPO₂ = 70%, the examination of the head and neck, cyanosis of the lips, and oral mucosa were observed. The baby had tachypnea and grunting. The large omphalocele was seen in the anterior part of chest and abdomen with 20 × 20 cm in diameter with a thin membrane. At the top of the omphalocele, a pulsatile mass was seen and in auscultation the heart’s grade III/VI murmur was heard. Abdominal viscera were visible inside the omphalocele. The heart was above, the liver in the middle and right, the spleen on the left, the stomach, small intestine, and the colon on the lower part of the masses. The baby was admitted to the intensive care unit. The site of the omphalocele was covered with sterile gas (mesh), and clinical/paraclinical measures were taken. Electrolytes and biochemical test were normal. In the arterial blood gas, there were metabolic acidosis and respiratory acidosis. The thoracoabdominal radiography reported a large omphalocele in the middle line, with the rise of the shadow of the liver and the absence of stomach air and left pulmonary edema. There was no pleural effusion (Fig. 1).

An echocardiography report ventricular septal defect, aortic arch displacement to right and pulmonary stenosis (Tetralogy of Fallot). Ultrasound reported a large omphalocele in the midline with the movement of the umbilical cord vessels upwards, the left lobe, and parts of the right lobe of the liver, a large part of the spleen on the left, the stomach, parts of the small intestine and colon were in the omphalocele. The gallbladder was not seen. There was no ascitis in the abdomen. Kidneys and bladder were normal. The heart was completely up and out of the chest. The kidneys and bladder were reported with normal size and echo. In the course of admission, due to respiratory failure, the baby needs intubation and mechanical ventilation. The baby died after 24 hours because of cardiopulmonary arrest, and unfortunately, the parents did not give autopsy consent (Fig. 2).

Discussion

The incidence of PC in most cases was sporadic and reported familial. Repeat on next pregnancy, associated with trisomy 13, 18, 21, and Turner’s syndrome were reported. Heredity attributed to chromosome X and gene mutation; thoracoabdominal syndrome (TAS) in the Xq25-q26.1 zone has been identified. The TAS gene plays a role in the sternum fusion, cardiac defects, and the abdominal wall aperture. New cases of bone morphogenic protein 2 (BMP2) gene mutations have been reported. This gene is responsible for the natural development of midline structures in the fetus. Additionally, mutations in the ALDH1A2 gene coded on chromosome 15 are reported to be responsible for the conversion of vitamin A to retinoic acid which plays a very important role in organogenesis during embryonic development.

The frequency of PC is 1 case per 1,000,000 live births and the ratio of male to the female is 2:1. In 1998, vazquez-jimenez et al, reported 153 cases of PC, 83% had heart abnormalities, 74.5% had abdominal wall defects, 60% had sternum defects, and 57% had diaphragmatic defects. The most common cardiac abnormality has been reported as ventricular septal defect.

The accompanying anomalies with PC consist of club foot, cleft lip and palate, spina bifida, meningocele, meningomyelocele, hydrocephalus, and encephalocele. The pathogenesis

Fig. 1 Simple thoracoabdominal radiography in newborn with PC. PC, Pentalogy of Cantrell.

Fig. 2 Anterior view of the infant with PC. PC, Pentalogy of Cantrell.
of the disease is unknown; however, the developmental defect
of the mesoderm layer on 14 to 18 days leads to the absence of
the formation of the horizontal diaphragm wall. This is fol-
lowed by the migration of visceral mesodermia over the
abdomen to the chest, the defective formation of sternum
and abdominal wall, and the inclusion of high visceral aper-
tures including the heart, parts of the pericardium which were
protrude into the cavity.
Some of the possible causes of pathogenesis include external
mechanical transduction to the embryo, external teratogenic
agents including the amniotic band resulting in tearing the yolk
sac in 14 to 18 days of the onset of conception, alcohol and drug
use, and maternal infections during pregnancy. Others include
viral infections in the first trimester of pregnancy and the use
of chlorine gas in pregnant women. The diagnosis of PC with the
help of ultrasound prior to the 12th week of conception is
impossible, since at this time, the abdominal viscera are outside
the abdominal cavity and in yolk sac. Prognosis of PC is poor and
varies depending on the severity of cardiac lesions and asso-
ciated anomalies.1 Deaths following surgery have been
reported in cases of cardiac abnormalities of up to 50%.1 The
main causes of mortalities are hypotension, shock, heart failure,
bradycardia, and cardiac arrest.

Conclusion
The presence of omphalocele in the fetus from 14th week of
pregnancy can be detected by ultrasound. Therefore, in all
cases with omphalocele, the presence of PC and associated
anomalies is recommended.

Compliance with Ethical Standards
Research involving human participants and/or animals—
ethical approval: All procedures performed in studies invol-
ving human participants were in accordance with the
ethical standards of the institutional and/or national
research committee and with the 1964 Helsinki declaration
and its later amendments or comparable ethical standards.

Informed Consent
Informed consent was obtained from all individual parti-
cipants included in the study.

Conflict of Interest
The authors deny any conflicts of interest in any terms or
by any means during the study. All the funds are provided
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