Twin reversed arterial perfusion sequence in a monochorionic monoamniotic twin pregnancy: a very rare condition

Abolfazl Abouie1, Nima Rakhshankhah1, Ladan Younesi2, Zeinab Safarpour2, Ayda Roostaee2 and Ghazaleh Amjad2*

Abstract

Background: Twin reversed arterial perfusion sequence (TRAP) is a very rare congenital anomaly. We present sonographic findings of TRAP sequence in the case of a multiparous woman with a monochorionic monoamniotic twin pregnancy who was referred to our unit for blood sugar control.

Case presentation: The patient had a history of co-twin demise at 13 weeks of gestation without appropriate fetal surveillance afterwards. We found a monochorionic placentation with a normal appearing pump twin, an abnormal appearing co-twin without obvious cardiac activity and reversed arterial flow toward instead of away from the anomalous acardiac fetus. Therefore, the sonographic diagnosis of TRAP sequence was confirmed.

Conclusions: We recommend considering the potential rare complications of monochorionic twin pregnancy which necessitates proper surveillance and intervention to monitor suitable growth of pump twin.

Keywords: Case report, TRAP sequence, Monochorionic monoamniotic, Twin pregnancy, Ultrasound

Background

TRAP sequence is a very rare congenital anomaly that occurs in 1/35,000 births [1]. This syndrome appears in nearly 1% of monochorionic twin pregnancy, with approximately 75% in diamniotic and 25% in monoamniotic twin pairs. Aberrant arterioarterial anastomosis in the single placenta accounts for specific complications in monochorionic twins [2]. In TRAP sequence, deoxygenated arterial blood pumps from pump twin to acardiac twin. Thus, in acardiac twin, lower structures have blood circulation and upper structures do not develop properly [3]. According to Malone classification [4], there are two kinds of twin in TRAP sequence which describe as hemicardius (imperfectly formed heart) and holocardius (absence of heart). According to Joseph C classification, acardiac fetus categorises into acardius anecps (partially developed head and face), acardius accephalus (no cephalic structures), acardius acormus (there is a head but no body) and acardius amorphous (no recognizable organs) [5]. In this case report, we present sonographic findings of a very rare case of TRAP sequence in monochorionic monoamniotic twin pregnancy.

Case presentation

A 22-year-old multiparous pregnant woman with gravida three, para one, living one and with a past history of one spontaneous abortion at 16 years of age was referred at 24 weeks of gestation to our tertiary obstetric and perinatology center for blood sugar control and fetal demise of one of the twins at 13 weeks of gestation. The patient had O+ blood group and medical history of hypothyroidism (consuming 50μg of levothyroxine daily) and gestational diabetes which had been controlled by dietary recommendations. The patient was asymptomatic at presentation, and laboratory studies...
such as blood count, coagulation status, biochemistry and urine analysis were all normal.

Abdominal ultrasonography showed intrauterine twin pregnancy with an anterior placenta without obvious membrane in favor of monochorionic monoamniotic twin pregnancy. The twin at the right side of uterine cavity represented cardiac activity and normal appearance, and the one at the left side was dysmorphic without detectable fetal heart activity (Fig. 1). The viable twin showed normal morphology and growth consistent with 24 weeks of gestation and normal fetal Doppler parameters (Fig. 2). In fetal echocardiography, no obvious structural cardiac anomaly was present. However, mild cardiomegaly and tricuspid regurgitation (TR) were evident. In addition, polyhydramnios with deepest fluid pocket (DFP) of 10.5 cm was evident of cardiac function failure (Fig. 3). On further examination of acardiac twin, a small and deformed skull with head circumference (HC) of 75 mm (equivalent to 13 weeks of gestation) was observed. Remarkable diffuse soft tissue edema was seen around the fetus with an average thickness of 33 mm, which was more severe in the lower limbs. The femoral length (FL) was 38 mm (compatible with 22 weeks of gestation) that represented lower extremity development in comparison with unrecognizable head, trunk and upper extremities (Fig. 1). Pulsed Doppler study of umbilical cord revealed reversed arterial flow toward rather than away from the anomalous fetus. Single umbilical artery also was obvious (Fig. 3). These ultrasound findings were all consistent with twin reversed arterial perfusion (TRAP) sequence in a monochorionic monoamniotic twin pregnancy. Three days

![Fig. 1](image1.png)

Fig. 1 a, b Dysmorphic acardiac fetus with diffuse soft tissue edema more severely in lower extremities. Marked growth discrepancy between HC and FL of acardiac twin is presented in c and d

![Fig. 2](image2.png)

Fig. 2 Normal fetal Doppler parameters such as umbilical artery (a), middle cerebral artery (b) and ductus venosus (c) in pump twin
later, the patient was referred with the complaint of rupture membrane, and cardiac activity was absent in both twins in ultrasonography.

Conclusions
The sonographic appearance of acardiac twin may differ in many cases, but a hydropic fetus without cardiac activity is the main manifestation of TRAP sequence. Reversed arterial flow in the umbilical artery from pump twin to acardiac twin is revealed in spectral Doppler examination [6]. Furthermore, most acardiac twin presents umbilical artery malformations, and according to Roberto Ruiz-Cordero, 78% of acardiac twin shows single umbilical artery and in 22% of cases, three vessel umbilical cord with thrombosis is described [7]. In majority of cases, pump twin develops high cardiac output failure including cardiomegaly, pericardial effusion and tricuspid regurgitation with polyhydramnios. Therefore, echocardiography is mandatory for the assessment and surveillance of cardiac function in pump twin [8]. The perinatal mortality of pump twin has been reported to range from 35 to 55% [9]. In our case, the pump twin demonstrated normal structural development. However, some degrees of cardiomegaly and mild tricuspid regurgitation in echocardiography and polyhydramnios were found, all of which were considered as a looming threat of fetal demise. All the abovementioned criteria associated with excessive volume of the acardiac twin predicted poor pregnancy outcome.

Intrauterine fetal demise and intra-amniotic or placental tumours such as placental teratoma should be considered as the differential diagnosis for TRAP sequence. However, precise evaluation of spinal development and umbilical cord attachment can be beneficial for differentiation [8].

There is a variety of treatment modalities which can improve perinatal survival. In minimally invasive techniques, vascular anastomosis is interrupted by alcohol, diathermy, cord embolization or coagulation, laser therapy or radiofrequency ablation [10]. Eventually, the most crucial component in treatment is likely to detect this complication early in pregnancy and perform appropriate interventions before detrimental consequences occur.

Abbreviations
TRAP sequence: Twin reversed arterial perfusion sequence; HC: Head circumference; FL: Femoral length

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Authors’ contributions
The authors have read and approved the final manuscript. All authors made a significant contribution to this study as mentioned below: AA: Data interpretation, writing the draft, discussion. NR: Data interpretation, writing the draft, discussion. LY: Study design, data collection, quality control. ZS: Study design, data collection, quality control. AR: Study design, data collection, quality control. GA: Study design, data interpretation, discussion.

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Availability of data and materials
The datasets used during the current report are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
The patient included in this study gave written informed consent to participate in this report.

Fig. 3 Single umbilical artery (a) with reversed flow to the acardiac fetus at pulsed Doppler study (b). Polyhydramnios (c) and mild cardiomegaly (d) in pump twin are signs of altered fetal heart function.
Consent for publication
Written informed consent to publish this information was obtained from study participant.

Competing interests
The authors have no conflict of interests.

Author details
1Department of Radiology, Iran University of Medical Sciences (IUMS), Tehran, Iran. 2Shahid Akbarabadi Clinical Research Development Unit (ShACRDU), Iran University of Medical Sciences (IUMS), Tehran, Iran.

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