Prenatal diagnosis of conjoined twins: four cases in a prenatal center

Yapışık ikizlerin prenatal tanısı: bir prenatal tanı üntesinin dört olgusu

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Abstract

Objective: To assess the findings in conjoined twins diagnosed prenatally.

Material and Methods: Between January 2002 and June 2009, we reviewed the database and medical records of 857 twin pregnancies, including 140 monochorionic twins. Nineteen monochorionic-monoamniotic twin pregnancies were detected, four of which were complicated by conjoined twins.

Results: Of these 4 cases, 2 were complicated by thoracopagus and one had thoraco-omphalopagus; these three cases underwent termination at 16, 11, and 19 weeks gestation, respectively. The last case was diagnosed as a pygopagus tetrapus parasitic twin at 28 weeks gestation. The family decided to continue the pregnancy, and achieved a successful outcome with elective surgery postpartum.

Conclusion: Conjoined twins are an uncommon and complex complication of monozygotic gestations, which is associated with high perinatal mortality. The early prenatal diagnosis of conjoined twins allows improved counseling about the management options, including maintenance of pregnancy with surgery after delivery or termination of pregnancy.

Key words: Conjoined twins, prenatal diagnosis

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Introduction

Conjoined twins are defined by the conjoined body area, and result from incomplete monozygotic twinning. The incidence of conjoined twins is estimated at 1% of monozygotic twinning or 1/200,000 live births (1). Over the last decade, the widespread use of assisted reproductive techniques has decreased the rate of monochorionic twins, and consequently may have changed the prevalence of conjoined twins (2, 3). Approximately 75% of conjoined twins are females (4). Conjoined twins have an anomaly of duplication in single-zygote conceptions which is thought to occur because of splitting and incomplete separation of the inner cell mass at 13-15 days post-fertilization. Forty percent of conjoined twins are stillborn and another 30% die during the 1st day of life (5). The prenatal diagnosis of conjoined twins began in 1976 (6). There is no doubt that the accurate and early diagnosis of fetal malformations will affect management and perinatal outcomes. We present four cases of conjoined twins (thoracopagus, thoracopagus, pygopagus tetrapus parasitic twins, and omphalopagus) detected in the first, second, and third trimesters using two-dimensional (2D) color Doppler ultrasound and three-dimensional (3D) ultrasound for the last case. Case 3 has previously been published with the diagnosis of a pygopagus tetrapus parasitic twin (7).

Material and Methods

All twin pregnancies admitted, followed, and delivered in our maternal-fetal medicine unit between January 2002 and June 2009 were recorded in medical charts and our computer database. A careful ultrasound evaluation was performed to identify or verify fetal details. Pregnancy and ultrasonographic findings were also entered into the database.

Twin pregnancies were divided into two groups: dichorionic (DC) and monochorionic (MC) pregnancies. MC...
twins were also sub-grouped as diamniotic and monoamniotic twins. Prenatal sonographic details were evaluated with related abnormalities depending on chorionicity and fetal development. Ultrasonographic examinations were performed transabdominally or transvaginally by one of four experienced maternal-fetal medicine physician sonographers using a Sonoline-G50 (TM-Siemens, multifrequency convex transducer 2.0-7.0 MHz; Issaquah, WA, USA) and Voluson 730 Expert (TM-GE Healthcare, multifrequency convex transducer 2.0-7.0 MHz, a 2-7-MHz convex transducer and a 4-8-MHz micro-4D convex real-time 4D-transducer; Milwaukee, WI, USA).

Results
During the study period, our database consisted of 857 twin pregnancies. We excluded 672 DC pregnancies with 140 MC twins. One hundred and two MC twins had regular development or additional malformations and 32 cases had twin-twin transfusion syndrome (TTTS). Nineteen cases were diagnosed as MC monoamniotic twin pregnancies, four of which were complicated by conjoined twins and two cases had acardiac twin development. Forty-five DC and MC twins were excluded because of an incomplete database.

Case 1
A 27-year-old gravida 1 para 0 was referred at 16 weeks 3 days gestation with a presumptive diagnosis of conjoined twins. On referral, the 2D transabdominal ultrasound demonstrated a monochorionic, monoamniotic twin gestation complicated by thoracopagus. Observations of the thoracic and abdominal cavities in transverse planes clearly demonstrated that the twins were joined over an area that extended from the central abdomen (omphalopagus) to the lower aspect of the thorax (thoracopagus). In view of the poor fetal prognosis, the family opted for pregnancy termination.

Case 2
A 19-year-old gravida 1 para 0 at 19 weeks gestation was registered for care in our prenatal clinic in the second trimester. Sonography revealed the presence of a monochorionic, monoamniotic twin gestation complicated by thoracopagus. Observations of the thoracic and abdominal cavities in transverse planes clearly demonstrated that the twins were joined over an area that extended from the central abdomen (omphalopagus) to the lower aspect of the thorax (thoracopagus). In view of the poor fetal prognosis, the family opted for pregnancy termination.

Case 3
A 30-year-old multigravida was referred to our maternal-fetal unit for evaluation of polyhydramnios at 28 weeks gestation. A sonographic examination revealed conjoined twins, each with a crown-rump length of 44 mm, consistent with 11 weeks 3 days gestation. The twins were joined at the abdomen. An omphalopagus conjoined twin pregnancy was diagnosed with 3D ultrasound (Figure 3). After counseling, the family opted for an elective abortion.

Discussion
Classically, conjoined twins have been reported to arise at around 12-13 days post-conception from a single blastocyst.
that had undergone incomplete division of the embryonic cell mass (4). Embryologic studies have shown that this extraordinary anomaly could develop from fusion of the two separate embryonic discs. Conjoined twins are classified according to the site of fusion and are always joined at identical anatomic points (Table 1). The combination of different types can occur, but the most frequently seen types include thoracopagus (20-40%), omphalopagus (18-33%), and parapagus (28%). Rachipagus is the rarest type of conjoined twins with an incidence of approximately 2% (8, 9).

The sonographic diagnosis of conjoined twins was first reported by Wilson et al. (6). Conjoined twins have been diagnosed in the first trimester with both transabdominal and transvaginal sonography (10, 11). According to Hill (12), the earliest prenatal suspicion of conjoined twins can arise at 7 weeks gestation. Because all conjoined twins are monochorionic and monoamnionic, the level of suspicion should increase when only one yolk sac is noted alongside two embryos in very early pregnancies or there are separating membranes not seen by ultrasound examination at any stage of pregnancy.

Non-operative management in cases with complex cardiac union, and emergency separation in cases with cardiac instability or additional structural anomalies warranting immediate surgical intervention and planned elective separation are ideally carried out between 2 and 4 months of age. Emergency separation carries a mortality rate of 71%, whereas the survival rate for elective separation is 80% (13). In cases with extensive cardiac or cerebral fusion, or when the anticipated severity of deformity following separation is unacceptable, the parents may desire to terminate the pregnancy (14). Although our cases with the diagnosis of separated hearts and thoracopagus and/or omphalopagus (cases 1, 2, and 4) had a better prognosis, the parents elected termination.

Although the prognosis for all types of conjoined twins is extremely poor, a careful anatomic and vascular mapping to determine the extent of organ sharing is of vital importance (15). First-trimester transvaginal ultrasonography in combination with color Doppler and 3D ultrasound are important advances that allow for the early diagnosis of conjoined twins. The data obtained by combining these complementary modalities may also be of prognostic value (16). Additionally, the accurate prenatal diagnosis of conjoined twins can be performed with these modalities as early as 10 weeks gestation (17, 18). Cases with false-positive diagnoses in the first trimester have been documented; therefore, an exact diagnosis should be made with caution (19). This technique may also provide images that are easier for parents to understand, which can help in decision making (20, 21).

Parasitic twins are a rare form of conjoined twins and consist of an incomplete twin (parasite) attached to the fully developed body of the co-twin (autosite). Parasitic twins are classified (22) as (a) an externally attached parasitic twin, (b) an enclosed fetus in fetu, (c) an internal teratoma, or (d) an acardiac twin connected via the placenta. This was the only surviving case (case 3) in our study in which the parents made the decision to continue the pregnancy and a successful result was achieved.

### Table 1. Classification of conjoined twins with incidence

<table>
<thead>
<tr>
<th>Classification</th>
<th>Incidence (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Ventral union</strong></td>
<td></td>
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<tr>
<td>Cephalopagus. Fusion from top of head to umbilicus. Each twin has 2 extremities and lower abdomen and pelvis are separated.</td>
<td>11</td>
</tr>
<tr>
<td>Thoracopagus. Twins are located face to face, with fused thoraces and shared heart or single interatrial vessel.</td>
<td>20-40</td>
</tr>
<tr>
<td>Omphalopagus. Twins have similar fusion to thoracopagus without shared heart or interatrial vessel.</td>
<td>18-33</td>
</tr>
<tr>
<td>Ischiopagus. These twins share a large conjoined pelvis and are more commonly joined end to end. External genitalia and anus are always shared.</td>
<td>6-11</td>
</tr>
<tr>
<td><strong>Dorsal union</strong></td>
<td></td>
</tr>
<tr>
<td>Craniopagus. Twins are joined by any portion of the skull except the face and foramen magnum. The bony cranium, meninges and brain are shared.</td>
<td>2</td>
</tr>
<tr>
<td>Pygopagus. Twins have fused sacrococcygeal and perineal regions, typically with shared anus but separate rectums. The spinal cord may be shared.</td>
<td>18-28</td>
</tr>
<tr>
<td>Rachipagus. Twins have dorsal fusion above the sacrum.</td>
<td>Rare</td>
</tr>
<tr>
<td><strong>Lateral union</strong></td>
<td></td>
</tr>
<tr>
<td>Parapagus. Twins have side-by-side connection with shared pelvis and variable cephalad sharing defined as follows:</td>
<td>28</td>
</tr>
<tr>
<td>- Dithoracic parapagus: separate thoraces and heads.</td>
<td></td>
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<tr>
<td>- Dicephalic parapagus: separate heads with fused thoraces.</td>
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<tr>
<td>- Diprosopus parapagus: 2 faces on the same side of single head.</td>
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</tbody>
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From Winkler et al. (9)
Conflict of interest
No conflict of interest is declared by authors.

References

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