US Evaluation of Twin Pregnancies: Importance of Chorionicity and Amnionicity

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Understanding Chorionicity and Amnionicty

Two separate fertilizations

Dizygotic pregnancy

1-3 days

Dichorionic Diamniotic (DiDi)

4-8 days

Monochorionic Diamniotic (MoDi)

8-13 days

Monochorionic Monoamniotic (MoMo)

13-18 days

Conjoined twins (MoMo)

Single fertilization (monozygotic)

Morula splits

Ultrasound in 1st trimester (7-9 weeks) can be up to 98% accurate in predicting chorionicity. Amnionicity can be challenging due to non-visualization of membranes. Repeat US, if necessary.
Why are Chorionicity and Amnionicity Important?

• Monochorionic (MC) twins account for 20% of twin pregnancies but 30% of all-cause pregnancy related complications.
• Single shared placenta hemodynamically connects two fetuses
• Rate of stillbirth in monochorionic compared to dichorionic twins:
  44.4 versus 12.2 per 1,000 births (relative risk: 3.6)
• Rate of neonatal mortality in monochorionic compared to dichorionic twins:
  32.4 versus 21.4 per 1,000 births (relative risk: 1.5)
• Twin pregnancies have a higher maternal risk of preterm labor, hypertensive disorders, diabetes, preterm labor, and premature rupture of membranes.
• Monochorionic diamniotic twins have unique complications including unequal placental sharing, twin-twin transfusion syndrome, twin anemia-polycythemia sequence, and twin reversed arterial perfusion syndrome.
• Monochorionic monoamniotic pregnancies have unique complications of conjoined twins, cord entanglement and higher rates of intrauterine demise.
• The later the age at which the morula splits, the higher the risk of congenital malformations.
Chorionicity and Amnionicity in the First Trimester

Number of Gestational Sacs = Chorionicity!
Number of Yolk Sacs = Amnionicity

Dichorionic Diamniotic twins:
Two thick chorionic sacs, each surrounding one embryo.

Monochorionic Diamniotic twins:
One thick chorionic sac surrounding both embryos. Each embryo surrounded by thin amnion. 2 Yolk sacs (Y).

Monochorionic Monoamniotic twins:
One thick echogenic chorion, one thin amnion, both embryos inside single amniotic sac.
Monochorionic Monoamniotic Pregnancy

Number of Yolk sacs = Amnionicity!

Single Yolk sac = Monoamniotic!

Embryos flanking each side of and sharing a single yolk sac = Monochorionic Monoamniotic pregnancy

Monochorionic twins: single yolk sac

Monochorionic twins: Single thick chorionic sac
Pitfall: Synechiae in Monochorionic Diamniotic Pregnancy Mimicking a Dichorionic Diamniotic Pregnancy

- **Suggestion of “thick” membrane**
  - 🚨 Same echogenicity as chorion

- **Other views demonstrate additional thin membranes??**

- **Single Chorionic Sac**

- **CONTINUOUS thin membrane, DISCONTINOUS thick membrane**

- **Chorionic reaction along synechiae**

- **Monochorionic Diamniotic pregnancy**
Pitfall: MoDi Pregnancy Mimicking a MoMo Pregnancy.
A case of missing membranes!

5 weeks 5 days

Single placenta = Monochorionic

11 weeks 3 days

Two embryos and two yolk sacs
No membrane present?
Diamniotic or Monoamniotic?

Follow up at 11w3d
Membrane now visualized

PITFALL: Membrane may be below sonographic detection in early MoDi pregnancy up to 7-9 weeks gestation

2 Yolk Sacs, one with each embryo

Embryo 1

Embryo 2

Thin membrane

Thin membrane
Single intrauterine pregnancy seen at REI with 2 mm crown-rump length, 8 week follow-up exam

Two for the price of one!

Single Yolk Sac

Single Amnion

Are they conjoined?

Difficult to be certain. Cardiac axis looks different. Follow-up needed to ensure embryos are separable.

Twin embryos present. Single Yolk Sac & Embryo = Monochorionic Monoamniotic
Chorionicity and Amnionicity in the Second Trimester

**Dichorionic twins:**
- Two placentas (separate or fused)
- Different gender = dizygotic
- THICK membrane
- Fused dichorionic placentas with TWIN PEAK SIGN

**Monochorionic twins:**
- Single placenta
- Same gender
- THIN inter-twin membrane

**Monoamniotic twins:**
- Single placenta
- Same gender
- No inter-twin membrane
- ENTANGLED CORDS

**Images:**
- **Different gender:**
  - GIRL
  - BOY
  - Thick membrane
- **Same gender:**
  - BOY
  - BOY
  - Thin membrane
  - Single placenta
- **No membrane:**
  - Entangled cords
  - Single placenta
Monochorionic Monoamniotic Conjoined Twins

Conjoined thoracopagus twins with shared hearts

- Incidence of conjoined twins: 10.25 per million births
- High mortality: Only 18% of prenatally diagnosed conjoined twins survive
- Increased incidence of structural anomalies not associated with conjoined organs

Thoracopagus

MRI redemonstrates conjoined twins with shared hearts

Specimen after pregnancy termination in a 4th case demonstrates thoraco-omphalopagus twins.
Complications Affecting a Monochorionic Pregnancy

Complications of unbalanced flow

1. Twin-twin transfusion syndrome (TTTS)
2. Twin anemia polycythemia sequence (TAPS)
3. Unequal placental sharing (*selective IUGR*)

Complications from in utero co-twin demise

1. High incidence of mortality.
2. Spectrum of mild to severe neurologic injury
3. Was called twin embolization; now known to be ischemic injury of survivor

Other complications

1. Twin reverse arterial perfusion syndrome (TRAP)
2. Discordant anomalies
3. Monoamniotic twins
4. Conjoined twins
Monochorionic Placenta & Placental Vascular Connections

Intraplacental vascular connections between the twin circulations

**Balanced State**
A-V inflow = outflow
A-A present

**Protective A-A Anastomosis**
 Twins pumping blood to each other

**Unbalanced State**
A-V inflow ≠ outflow
Blood flow from one twin to another

Normal vascular supply to single twin, no communication to other twin
Effects of Unbalanced Flow in a Monochorionic Twin Pair

1. Anemia
2. Oliguria
3. Oligohydramnios
4. Circulatory insufficiency
5. Growth restriction
6. Heart failure
7. Fetal demise

Donor twin often has marginal or velamentous placental cord insertion

Disease states:
1. Twin-twin transfusion syndrome (TTTS)
2. Twin anemia polycythemia sequence (TAPS)
3. Unequal placental sharing

Unbalanced shunt flow

Donor

Recipient

Placenta
Surveillance of Monochorionic Twins: Initiation, Timing, and Duration

- Initial scan—confirm living monochorionic pair
- Establish twin identity at first scan—identify placental cord insertion, discordant anomalies, or growth
- **Maintain identity for all future follow-ups**
- Ultrasound every 2 weeks → Starting at 16 weeks to delivery
- Growth scan q 4 weeks per NAFTA recommendations (*institutional protocols vary from 2-4 weeks*)

**TWIN-TWIN TRANSFUSION SYNDROME**
- Bladder and amniotic fluid
- Multivessel Doppler- Umbilical Artery, Vein and Ductus venosus

**TWIN ANEMIA POLYCYTHEMIA SEQUENCE**
- Middle cerebral artery velocity measurement
- Calculate Multiples of Median (*ref: perinatology.com*)

**UNEQUAL PLACENTAL SHARING/SELECTIVE IUGR**
- Check fetal weights
- Calculate weight discordance

- Anatomy scan at 18-22 weeks
- **Role of Fetal Echocardiography**—Risk of cardiac defects in monochorionic twins, both embryologic defects and physiologic changes from unique placental vascular connections
  - Perform at 18-22 weeks in uncomplicated monochorionic twin gestation
Monochorionic Diamniotic Pregnancy with Unequal Placental Sharing

- Single anterior placenta shared by monochorionic twins
- Weight discordance 30%
- Velamentous cord insertion of the smaller twin

Calculating weight discordance

Larger twin weight - Smaller twin weight

>25% weight discordance and/or fetal weight < 10th percentile constitutes unequal placental sharing
MoDi Pregnancy with Unequal Placental Sharing

- Normal UA Doppler with forward diastolic flow (yellow arrows)
- Normal antegrade umbilical vein flow without pulsatility (white arrow)
- Occasional loss of end-diastolic flow can sometimes be seen in the setting of an A-A (blue arrows)

- Arterial-arterial (A-A) anastomosis is demonstrated with *to and fro* flow
- Sometimes protective against TTTS
- Creates a balanced state
- To look for an A-A, search along the placental surface with color Doppler
**Stage 1 Twin-Twin Transfusion Syndrome**

**Quintero staging for TTTS**

- **Stage I**: Oligohydramnios in donor twin and polyhydramnios in recipient twin (Oli-Poly)
- **Stage II**: Absent bladder in the donor twin
- **Stage III**: Doppler abnormalities in the umbilical artery and/or umbilical vein
- **Stage IV**: Fetal hydrops
- **Stage V**: Fetal demise

**Oli-Poly**
- Absent bladder
- Doppler abnormalities
- Fetal hydrops
- Fetal demise

1. Prognostic value
2. Stepwise progression is expected but not always the norm
3. Often twin pairs can “skip” stages with progressive TTTS
Stage 2 TTTS Complicating a Monochorionic Diamniotic Pregnancy

- **Polyhydramnios (Donor)**: DVP = 9.8 cm
- **Oligohydramnios (Recipient)**: DVP = 0.7 cm
- **Bladder present (Recipient)**
- **Donor oligohydramnios** & **Recipient polyhydramnios**
- **Donor bladder non-visualized**
- **Bilateral umbilical arteries expected to encircle the bladder**

**Single placenta**

**Stuck twin (donor)**

**Note difference in echogenicity of the 2 amniotic cavities**

**Stage 2 TTTS**
Stage 2 TTTS Complicating a Monochorionic Monoamniotic Pregnancy

1. Challenging diagnosis due to single amniotic space.
2. Polyhydramnios is the clue.
3. Absent bladder in one of the twins = Stage 2 TTTS
Stage 3 Twin-Twin Transfusion Syndrome

Stuck twin (donor), anhydramnios

- Bladder absent in donor
- Polyhydramnios
  - DVP = 9.5 cm in recipient

Planning images demonstrate sonographic window for intraoperative guidance of laser ablation procedure.

- Absent end diastolic flow in donor
- Donor oligohydramnios & Recipient polyhydramnios
- Donor absent bladder
- Doppler abnormality in Donor

= Stage 3 TTTS
Stage 4 Twin-Twin Transfusion Syndrome

**Donor**

- Stuck Donor Twin with *anhydramnios*
- Recipient twin with polyhydramnios and *hydrops* (ascites shown here)

**Recipient**

- Recipient umbilical artery waveform shows *absent end diastolic flow* (arrows)
- Recipient umbilical vein waveform shows *pulsatile flow* (arrows). *Pulsatile flow in the UV is a premorbid waveform*

**Stage IV TTTS**
Treatment for Twin-Twin Transfusion Syndrome

- **Fetoscopic laser ablation**
  - Quintero stage II–IV disease
  - Gestational age between 16 weeks 0 day and 26 weeks 0 day of gestation
  - Maternal complications such as placental abruption (1%) and intra-abdominal leakage of amniotic fluid (3%)
  - Goal: Laser coagulation of the entire vascular equator

- **Ultrasound surveillance with MCA Doppler after treatment**
  - Weekly to assess for recurrent twin-twin transfusion syndrome or the development of twin anemia–polycythemia sequence for 6 weeks
  - Then, every 2 weeks as with uncomplicated monochorionic gestation

- **Amnioreduction** for non-laser candidates (after 26 weeks)

- **Selective feticide/Radiofrequency ablation** (rarely performed):
  - Usually performed after 16 weeks
  - Discordant for a major structural abnormality
  - One of the twins is moribund
  - Severe IUGR of one twin
Complication after Laser: Chorioamniotic Separation

Post procedurally, a membrane is seen outlining the amniotic surface of the placenta and continuing along the uterine body, representing the amnion separated from the chorion. Shredded membranes were also visible.

**Understanding the Chorioamniotic space:**
The chorion and amnion are separate in early pregnancy and fluid is present between them. Eventually, this potential space is obliterated and fluid can re-enter this space as a procedural complication. **FACT:** Chorion and amnion never fuse with each other; this is a common misconception.

**Correct method of measuring DVP after CAS.**
Space outside the amnion should *not* be included in the measurement.
Twin Reverse Arterial Perfusion Sequence

28 year old woman with twin pregnancy, provided history of embryonic demise of one twin

Single placenta/chorion

“Demised” twin

Living twin

Follow-up after 6 weeks

Reversed umbilical arterial flow in anomalous twin

Interval growth of “demised” twin

Interval growth of anomalous embryo

severely edematous fetus

Reversed umbilical arterial flow

= Twin Reverse Arterial Perfusion Sequence
Twin Reverse Arterial Perfusion Sequence

- Monochorionic twins
- Abnormal placental connections
- TRAP twin abnormal head, torso
  - May have NO cranial structures
- Can have a rudimentary heart
  - Acardiac twin is a misnomer

Outcomes:
1. Self devascularization
2. Pump twin hydrops/demise
   (~50% chance of pump twin demise)
3. Treat with radiofrequency ablation

• Upper part of fetus fails to develop, no well developed cranial structures
• Can have a rudimentary heart

Placenta
Abnormal flow to TRAP through A-a connections

Deoxgenated blood pumped to the TRAP fetus. Pump twin at risk of high output failure
TVUS at 10 weeks showed monochorionic diamniotic twins. Both demonstrated cardiac activity. One was hydropic and lacked normal cranial structures. The other was normal.

TVUS at 12 weeks showed demise of both the reverse perfused twin and the pump twin.

The abnormal twin cannot survive ex-utero and management of this condition is focused on wellbeing of the pump twin. Hence, radiofrequency ablation of the abnormal twin is performed.
Radiofrequency Ablation of Abnormal TRAP Twin

Abnormal twin with perfusion and flow within the cord (arrow) demonstrated on color Doppler US.

Spectral Doppler US demonstrates *arterial flow flowing towards the abnormal twin*, consistent with twin reverse arterial perfusion sequence.

Correct placement of a radiofrequency ablation (RFA) device with the *tip at the abdominal cord insertion*. Unlike the fetoscope, the RFA device can be placed through the placenta due to its smaller size.

Post-delivery image demonstrating the anterior abdominal wall defect post RFA.

No residual flow within the TRAP twin after RFA.
Pitfall: Post RFA of the Abnormal TRAP Twin

Note the high scale for Doppler interrogation. Such a high scale coupled with high gain settings predisposes to twinkle artifact.

32 week MoDi pregnancy after RFA of TRAP twin. Color Doppler images demonstrate a large amount of signal throughout the TRAP twin. This represents “twinkle artifact” from the osseous structures of the abnormal twin.

**PITFALL:** Do not misinterpret as presence of flow.

Adjusting the scale lower and using appropriate gain can help reduce artifact and assess true flow to the TRAP twin.
**Twin Anemia Polycythemia Sequence**

- Gestational age: 19 wks 3 days
- Normal fluid in both sacs (*)

**Calculating Multiples of Median**

**Discordant MCA PSV**

| + | Normal fluid and bladders |

**Twin Anemia Polycythemia Sequence**

**Interpreting Multiples of Median**

- **Polycythemia:** <1.0 MoM
- **Anemia:** >1.5 MoM

**Expected Peak Velocity of Systolic Blood Flow in the MCA as a Function of Gestational Age**

Enter values here:

- Gestational age (weeks)
- Observed MCA Peak Systolic Velocity (cm/sec)

Calculations: The Median Peak Systolic Velocity for this age is [ ] MoM

Your measurement is [ ] MoM

Results
Demise of One Twin of a Monochorionic Pair

- Subsequent high risk of death & severe cerebral injury to surviving twin
- Suspected mechanism: *Acute exsanguination* of the surviving twin into the *low-pressure vascular circuit of the deceased twin* through patent vascular anastomoses
- → Sudden and profound hypotension, hypovolemia, and anemia
- → → Consequent *tissue hypoxia and acidosis* (earlier theory of intertwin embolization has largely been abandoned)
- Urgent delivery after an unwitnessed twin death unlikely to improve the co-twin’s outcome and may unnecessarily expose the survivor to complications of prematurity.
- *Expectant management* to term is favored

Porencephaly with anechoic spaces lining the ventricular wall, in a previously normal brain after co-twin demise

Dilated heart in a surviving twin after co-twin demise, representing ischemic cardiomyopathy in the surviving twin. This twin also had ischemic brain injury.
Demise of one twin of a monochorionic pair

Surviving twin demonstrates large right parietal infarct and diffuse left cerebral hemisphere ischemia. The brain was previously structurally normal.

Corresponding fetal MR images (a) T2-weighted (b) diffusion and (c) ADC sequences: demonstrate large right parieto-temporal infarction with large area of cortical necrosis and diffuse left cerebral hemisphere ischemia.

Ischemic injury is possible to multiple organ systems

- **CENTRAL NERVOUS SYSTEM**
  - Ventriculomegaly
  - Porencephaly
  - Cerebral atrophy
  - Microcephaly

- **CARDIOVASCULAR SYSTEM**
  - Ischemic cardiomyopathy

- **GASTROINTESTINAL TRACT**
  - Small bowel atresia

- **GENITOURINARY TRACT**
  - Renal cortical necrosis
Discordant Anomalies

- Monochorionic twins assumed to be genotypically identical
- May be *phenotypically discordant for major congenital malformations*
- Structural anomalies more common than in singletons & dichorionic twins
- Major anomalies: 6–8% of monochorionic twins vs 1–2% of dichorionic twins
- *Usually affects only one fetus*
- Discordant anomalies *may also negatively affect healthy co-twin*
- Intrauterine demise due to anomalies or aneuploidy \(\rightarrow\) Co-twin at risk of demise (10–25%) or cerebral damage (24–45%)
Higher-Order Multiples

- High-order multifetal gestations containing a monochorionic twin pair are at increased risk of adverse pregnancy outcomes.
- Lower birth weight
- Lower gestational age at delivery
- Spend more days in the NICU
- Increased risk of intrauterine death, selective growth restriction, delivery less than 32 weeks of gestation, and a lower survival rate in dichorionic triplets than in trichorionic triplets.
- Option of reducing the monochorionic pair.
Referral to a Fetal Treatment Center

Monochorionic pregnancy

- No
- Yes

Multiple gestation protocol

- No
- Yes

Gestational age

- > 34 weeks
- < 34 weeks

Delivery

Oligohydramnios/Polyhydramnios

- No
- Yes

Twin twin transfusion syndrome

Additional testing and referral to a fetal treatment center

Weight discordance >25%

- No
- Yes

MCA discordance

Twin Anemia polycythemia sequence

Gestational age

- > 34 weeks
- < 34 weeks

Delivery

Oligohydramnios/Polyhydramnios

- No
- Yes

Twin twin transfusion syndrome

Additional testing and referral to a fetal treatment center

Establish chorionicity as soon as possible!!
Thank you for viewing our presentation!

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References:


