

Delayed Interval Delivery of the Second Twin: Neonatal and maternal outcomes



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Introduction

In Spain, due to the increase of assisted reproductive techniques, the multiple pregnancy rate level up to 2.06% vs 1% around the world¹.

More than 20% of twins are born preterm². Prematurity is the main cause of morbility and mortality in children <5 years and it is inversely correlated to the gestational age of preterm birth³.

Delayed interval delivery (DID) of the second fetus is a management option in some selected cases to prolong gestational age of the remaining fetus, in order to improve survival and neonatal outcomes^{4,5}.

Objective

To evaluate neonatal and maternal outcomes of the remaining fetus after DID in twin pregnancies.



Retrospective observational study between 2014-2022

- Twin DC or MCDA
- Delivery <28 weeks without preterm premature rupture of membranes (pPROM), haemorrhage, infection and fetal wellbing of the remaining fetus.

After deliverv of first fetus

- Umbilical cord ligated. Placenta was left inside the uterus.
- Amniocentesis of the remaining amniotic sac to rule out intra-amniotic infection
- Antenatal management:
- Broad-spectrum antibiotic prophylaxis
- Fetal lung maturation (betamethasone) after X 23+0 weeks
- Fetal neuroprotection (magnesium sulphate) if imminent delivery was suspected before 32+0 weeks
- Tocolytics were only given if uterine contractions after exclusion of clinical chorioamnionitis.

Cerclage was not placed after DID.

Table 1: Basal characteristics and delivery

Results

14 cases were included.

Latency from delivery of the first twin to delivery of the second twin was of 9 (2.5; 37.5) days.

Neonatal survival rate to discharge home of the second twin was 64.3%. Despite not being statistically significant a lower neonatal morbidity composite rate was observed in the second twin.

Maternal composite morbidity rate was 17.9%. No cases of maternal sepsis, admissions to ICU or hysterectomy were observed.

Conclusion

DID in selected extreme preterm birth twins increase gestational age at delivery and, therefore, birthweight of the remaining fetus without increasing neonatal morbidity outcome and with a low maternal morbidity associated.

Maternal age (years)	35.5 (32.0-39.2)
Body mass index (kg/m2)	24.7 (28.3-30.4)
Nulliparous	10 (71.4)
Pregnancy after In vitro fertilization treatment	6 (42.9)
Bichorial biamniotic pregnancy	13 (92.9)
Monochorial biamniotic pregnancy	1 (7.1)
Cerclage prior to DID	4 (28.5)
Sestational age at admission (weeks)	23.6 (19.5-25.6)
Cervical lenght at admission (mm)	3.5 (0-25.5)
Positive vaginal culture at admission	1 (7.1)

Table 18.7: Continuous variables were presented as medians	(25th, 75th
Maternal composite morbidity	5 (17.9)
Postpartum hemorrhage treated with drugs different than oxytocin	2 (15.4)
Postpartum endometritis	1 (7.7)
Maternal clinical chorioamnionitis	0

percentile).

Table 3. Neonatal outcomes

	First twin	Second twin	p value
WBC at delivery	11655 (10400- 16075)	12545 (10225- 17462)	0.550
CPR at delivery (mg/L)	12.5 (4.5-17.5)	57 (50-65)	0.034
Gestational age at delivery (weeks)	23.6 (19.1-26.3)	28.2 (25.0-30.0)	0.018
Vaginal delivery	14 (100)	10 (71.4)	0.031
Emergency cesarean section	0	2 (14.3)	0.277
Female gender	5 (45.5)	7 (53.8)	0.682
Birth weight (g)	515 (420-714)	1100 (570-1215)	0.013
Apgar at 1 minute <7	9 (90)	4 (34.6)	0.011
Neonatal survival rate to discharge home	6 (42.9)	9 (64.3)	0.376*
Stillbirth	4 (40)	1 (12.5)	0.220*
Neonatal death	4 (57.1)	2 (25)	0.192*
Respiratory distress syndrome or transient neonatal tachypnea	7 (87.5)	6 (85.7)	0.989*
Bronchopulmonary dysplasia	6 (75)	2 (28.6)	0.427*
Intraventricular hemorrhage grade III or IV	2 (25)	3 (42)	0.480*
Necrotizing enterocolitis	1 (12.5)	1 (14.3)	0.071*
Early onset sepsis with positive	4 (50)	1 (14.3)	0.167*

Continuous variables were compared using a nonparametric U Mann-Whitney test presented as medians (25th; 75th percentile). Categorical variables were compared using Chi-square or

Fisher exact tests and presented as number (%)

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Primary outcome measures were

composite outcomes for maternal

morbidity (Table 2) and neonatal

morbimortality (Table 3)