Effect of birth order on neonatal morbidity and mortality among very low birthweight twins: a population based study

E S Shinwell, I Blickstein, A Lusky, B Reichman


Objective: To study the effect of birth order on the risk for respiratory distress syndrome (RDS), chronic lung disease (CLD), adverse neurological findings, and death in very low birthweight (VLBW; < 1500 g) twins.

Methods: A population based study of VLBW infants from the Israel National VLBW Infant Database. The sample included all complete sets of VLBW twin pairs admitted to all 28 neonatal intensive care units between 1995 and 1999. Outcome variables were compared by birth order and stratified by mode of delivery and gestational age, using General Estimating Equation models, with results expressed as odds ratio (OR) with 95% confidence interval (CI).

Results: Second twins were at increased risk for RDS (OR 1.51, 95% CI 1.29 to 1.76), CLD (OR 1.36, 95% CI 1.11 to 1.66), and death (OR 1.24, 95% CI 1.02 to 1.51) but not for adverse neurological findings (OR 1.20, 95% CI 0.91 to 1.60). Mode of delivery did not significantly influence outcome. The odds ratio for RDS in the second twin was inversely related to gestational age, and the increased risk for RDS and CLD was found in both vaginal and caesarean deliveries.

Conclusions: VLBW second twins are at increased risk for acute and chronic lung disease and neonatal mortality, irrespective of mode of delivery.

P reterm second twins are at increased risk for respiratory distress syndrome (RDS). Although this observation has been recognised for many years, improved obstetric and neonatal care, including antenatal steroids and postnatal surfactant, have improved the outcome of high risk very low birthweight (VLBW; < 1500 g) infants, and therefore this observation may no longer be valid. Moreover, previous reports have mostly studied relatively small series of infants.

Accordingly, we evaluated a large population database for the effect of birth order on RDS, chronic lung disease (CLD), adverse neurological findings, and mortality in VLBW twins. As previous studies have related the increased risk of second twins to malpresentation associated asphyxia, this study also examines the interaction of perinatal depression, malpresentation, and gestational age with the effect of birth order on these outcomes.

METHODS

Israel National VLBW Infant Database

Infants with birth weight of 1500 g or less who are born alive in all of the country’s 28 neonatal units are included in the database. Data are prospectively collected on a prestructured form and include information on the parents, maternal pregnancy history and antenatal care, mode of delivery, infant’s status at birth, procedures and morbidity during hospital stay, and outcome at discharge. Definitions used are concomitant with those of the Vermont-Oxford Trials Network and were defined by the scientific committee before data collection and have remained unaltered since. Once collected by the local investigators, the data are sent to the database coordinator, checked for missing items and logic errors, corrected, completed, and then entered into a computerised database. Patient information is cross checked with the national birth registry, and any missing data are requested from the birth hospital. Data are collected on all infants until discharge home or death.

Study sample

For the purposes of this study, we included all sets of twins in which both twins had a birth weight of 1500 g or less and gestational age of 24 weeks or more. The vast majority (86%) were of gestational age 32 weeks or less. Sets in which one member did not meet the criteria were excluded. We also excluded 31 cases of caesarean section for the second twin after vaginal delivery of the first twin.

From 1995 through 1999, the database included records of 7047 infants which comprise 99% of all VLBW live births in Israel (approximately 1.2% of all births). Thirty percent (2131) of the infants were twins, and 1328 infants (664 twin pairs) fulfilled the study criteria. In the subsequent analysis of major neonatal morbidities, further exclusions were due to neonatal deaths of one or both members of the twin pair, or missing data. The analysis of RDS included 651 pairs after exclusion of 13 delivery room deaths. CLD was evaluated in 434 pairs, excluding neonatal deaths or missing data (n = 24) related to oxygen therapy at 28 days of age. Adverse neurological findings were considered in 412 pairs, excluding twin pairs in whom one or both infants died or an appropriate ultrasound was not performed (n = 46). The minimum requirement for ultrasound was two examinations, one before age 28 days for identification of intraventricular haemorrhage and one after age 28 days for periventricular leucomalacia and ventricular dilatation.

Definition of outcome variables

Death was defined as occurring during the hospital stay. Delivery room resuscitation included endotracheal intubation and/or cardiac massage or adrenaline (epinephrine) administration. RDS was defined by characteristic clinical and radiographic findings. CLD was defined as a clinical diagnosis

Abbreviations: CLD, chronic lung disease; RDS, respiratory distress syndrome; VLBW, very low birthweight
Statistical analysis
For each outcome variable, we used the Generalised Estimating Equation approach to estimate and test the odds ratios for first versus second twins. In a univariate analysis, the odds ratio estimate using this method is equal to the usual estimate of odds of the outcome among second twins divided by the estimate of odds of outcome among first twins. We chose this estimate as it included the data from all sets of twins and not only the discordant sets. The Generalised Estimating Equation method takes into account the correlation between the outcomes of sibling pairs.

To test if the odds for second versus first twins ratios were different for twins delivered vaginally compared with caesarean section and by groups of gestational age, stratified analysis was performed by mode of delivery and gestational age. Results are presented as odds ratio (OR) and 95% confidence interval (CI). Analyses were performed using the SAS (SAS Institute, Inc, Cary, North Carolina, USA) statistical software (version 8.2).

RESULTS
Characterisation of study sample
The study population comprised 664 VLBW twin pairs. The mean (SD) gestational age at delivery was 28.4 (2.4) weeks, and the birth weights of the first and second twins were similar (1072 (274) v 1056 (270) g). Antenatal steroids were administered in 70%, and 65% were delivered by caesarean section.

Outcome variables
The incidence of the outcome variables in the study sample was RDS 70%, CLD 19%, adverse neurological findings 39%, and mortality 21%. Table 1 shows the comparison of the outcome variables between the first and second twins. Second twins were at significantly higher risk for RDS (OR 1.51, 95% CI 1.29 to 1.76), CLD (OR 1.36, 95% CI 1.11 to 1.66), and death (OR 1.24, 95% CI 1.02 to 1.51). No significant difference was found between the groups of twins in the risk for adverse neurological findings (OR 1.20, 95% CI 0.91 to 1.60). Most twin pairs were concordant for each of the outcome variables (death 81%, RDS 82%, CLD 88%, adverse neurological findings 79%).

Table 2 shows similar odds ratios for each of the outcomes in twin pairs born vaginally or by caesarean section. The excess risk in second twins for RDS and CLD was significant in infants delivered by caesarean section or vaginally. However, as the sample in this stratified analysis was considerably smaller, the odds ratio for mortality in these subgroups did not quite meet statistical significance.

<table>
<thead>
<tr>
<th>Variable</th>
<th>No of twin pairs</th>
<th>First twin (%)</th>
<th>Second twin (%)</th>
<th>OR</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Death</td>
<td>664</td>
<td>19.6</td>
<td>23.2</td>
<td>1.24</td>
<td>1.02 to 1.51</td>
</tr>
<tr>
<td>RDS</td>
<td>651</td>
<td>65.7</td>
<td>74.3</td>
<td>1.51</td>
<td>1.29 to 1.76</td>
</tr>
<tr>
<td>CLD</td>
<td>434</td>
<td>18.4</td>
<td>23.5</td>
<td>1.36</td>
<td>1.11 to 1.66</td>
</tr>
<tr>
<td>Neuro</td>
<td>412</td>
<td>18.2</td>
<td>21.1</td>
<td>1.20</td>
<td>0.91 to 1.60</td>
</tr>
</tbody>
</table>

RDS, Respiratory distress syndrome; CLD, chronic lung disease; Neuro, adverse neurological findings; OR, odds ratio; 95% CI, 95% confidence interval.

Perinatal status
Second twins were at higher risk for malpresentation (OR 2.10, 95% CI 1.71 to 2.57). However, no difference was found between the twin groups in Apgar scores at five minutes (OR 1.05, 95% CI 0.86 to 1.28) or in the need for resuscitative measures in the delivery room (OR 1.08, 95% CI 0.95 to 1.22).

Gestational age
To assess the risk for RDS and CLD while controlling for gestational age, we calculated odds ratios for these outcome variables in subgroups of 24–27 weeks, 28–30 weeks, and greater than 30 weeks (table 3). The increased risk for RDS in second twins appeared to be inversely related to gestational age, with maximum effect in the 24–27 week group. The increased risk for CLD was found to be significant only in the 24–27 week gestation group. The other outcomes were not compared by gestational age because of low incidence.

DISCUSSION
This study confirms that VLBW second twins are at significantly increased risk for RDS, CLD, and death and that the excess risk for RDS and CLD is most pronounced in infants born at 24–27 weeks gestation. In addition, it is of note that delivery by caesarean section does not appear to protect the second twin from this increased risk.

This study has certain strengths and weaknesses that may influence the interpretation of the findings. The major strength is the large, population based sample of high risk VLBW neonates, drawn from both local general and university teaching hospitals. The focus on VLBW infants increased the likelihood of finding real differences between the twin groups, and further sensitivity was achieved by the use of within pairs statistical analyses compared with the non-paired approach used in the studies of Chen et al and Neilsen et al.

However, a number of limitations need to be discussed. Firstly, although the sample was large, it was not large enough to resolve the issue of the influence of gestational age on the increased risk for RDS in second twins. Hacking et al found a direct correlation with gestational age, whereas this study, with a larger sample, found an inverse correlation. To study this question more satisfactorily, a much larger sample will be required. Another concern is the definition of CLD as a clinical diagnosis of bronchopulmonary dysplasia and requirement for supplemental oxygen therapy at 28 days of life. The studies of Hacking et al and Donovan et al both found second twins to be at increased risk for RDS but not for CLD, when defined as oxygen requirement at 36 weeks gestational age. The shorter term definition was used in our study because of the higher availability of the information in the database. Clearly, as the incidence of oxygen dependency is lower at 36 weeks, there is less likelihood of finding significant intertwin differences or, alternatively, there may be a real difference at 28 days which is diminished by 36 weeks. A larger study with reliable data at both time points will be required to resolve this.
The influence of birth order on the other outcome variables remains less clear. We found a small but significant increase in mortality risk (OR 1.24, 95% CI 1.02 to 1.51), which was not found in the studies of VLBW infants and of all preterm infants. None of the population based studies found an increase in intraventricular haemorrhage or adverse neurological findings.

Previous studies have suggested that the increased risk for RDS in non-presenting twins is only found in infants delivered vaginally and that this relates to the increased incidence of malpresentation and presumed “asphyxia” in the second twin. This and other studies did not confirm such findings. Although second twins had increased risk for malpresentation, there was no increased risk for low five minute Apgar scores or need for resuscitative measures. It follows that clinically evident asphyxia is not part of the cause of the increased risk of second twins for RDS. However, an untested possibility is that transient hypoxia and/or acidosis may affect surfactant function, which may contribute to the development of mild respiratory distress.

In summary, we conclude that, even in the era of widespread use of antenatal steroids, postnatal surfactant, and modern intensive care, VLBW second twins appear to have an increased risk for acute and chronic lung disease and death. The mechanisms of this effect remain partially understood.

ACKNOWLEDGEMENTS

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REFERENCES


Table 2  Odds ratios and 95% confidence intervals for outcome variables in first and second twins delivered vaginally or by caesarean section

<table>
<thead>
<tr>
<th>Vaginal delivery</th>
<th>Coaearane section</th>
</tr>
</thead>
<tbody>
<tr>
<td>No of twin pairs</td>
<td>OR (95% CI)</td>
</tr>
<tr>
<td>Death</td>
<td>233</td>
</tr>
<tr>
<td>RDS</td>
<td>227</td>
</tr>
<tr>
<td>CLD</td>
<td>142</td>
</tr>
<tr>
<td>Neuro</td>
<td>131</td>
</tr>
</tbody>
</table>

RDS, Respiratory distress syndrome; CLD, chronic lung disease; Neuro, adverse neurological findings; OR, odds ratio; 95% CI, 95% confidence interval.

Table 3  Effect of gestational age on odds ratios for respiratory distress syndrome (RDS) and chronic lung disease (CLD) in first and second twins

<table>
<thead>
<tr>
<th>GA (weeks)</th>
<th>n</th>
<th>OR (95% CI)</th>
<th>n</th>
<th>OR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>24–27</td>
<td>216</td>
<td>2.48</td>
<td>1.31 to 4.69</td>
<td>84</td>
</tr>
<tr>
<td>28–30</td>
<td>304</td>
<td>1.71</td>
<td>1.34 to 2.19</td>
<td>233</td>
</tr>
<tr>
<td>31+</td>
<td>131</td>
<td>1.42</td>
<td>0.95 to 2.12</td>
<td>117</td>
</tr>
</tbody>
</table>

GA, Gestational age; n, number of twin pairs; OR, odds ratio; 95% CI, 95% confidence interval.


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