Changes in pulmonary arterial pressure in preterm infants with chronic lung disease

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Abstract

Background—Pulmonary arterial pressure (PAP) is raised in preterm infants with respiratory distress syndrome who subsequently develop chronic lung disease. The natural history of pulmonary hypertension in infants with chronic lung disease is unknown.

Objectives—To investigate changes in PAP, assessed non-invasively using Doppler echocardiography, in infants with chronic lung disease during the 1st year of life.

Methods—Serial examinations were performed in infants with chronic lung disease and healthy preterm infants. The Doppler derived acceleration time to right ventricular ejection time ratio (AT/RVET) was calculated from measurements made from the pulmonary artery velocity waveform.

Results—A total of 248 examinations were performed in 54 infants with chronic lung disease and 44 healthy preterm infants. The median AT/RVET was significantly lower in infants with chronic lung disease than in healthy preterm infants (0.31 versus 0.37). AT/RVET significantly correlated with age corrected for prematurity in both infants with chronic lung disease (r = 0.67) and healthy infants (r = 0.55). There was no significant difference between the rate of change in AT/RVET between the two groups. In infants with chronic lung disease, multivariate analysis showed that AT/RVET was significantly independently associated with age and inversely with duration of supplemental oxygen treatment. Median AT/RVET was significantly lower in infants with chronic lung disease until 40–52 weeks of age corrected for prematurity.

Conclusions—Although PAP falls with increasing age in both infants with chronic lung disease and healthy preterm infants, it remains persistently raised in infants with chronic lung disease until the end of the 1st year of life. (Arch Dis Child Fetal Neonatal Ed 2000;82:F243–F247)

Keywords: pulmonary arterial pressure; chronic lung disease; Doppler echocardiography; acceleration time to right ventricular ejection time ratio; prematurity

Since Northway et al first described the pathological features of bronchopulmonary dysplasia, a wide spectrum of cardiorespiratory abnormalities has been recognised. In particular, several abnormalities of structure and function of the pulmonary circulation have been reported.2,4

Pulmonary arterial pressure (PAP) is initially raised in infants with respiratory distress syndrome and, in most, gradually declines with improvement in respiratory status.5,9 However, PAP remains persistently raised in infants who subsequently develop chronic lung disease.8,10 Pulmonary hypertension with raised pulmonary vascular resistance in infants and older children with severe established chronic lung disease has also been demonstrated at cardiac catheterisation in a number of studies, and is associated with a poor prognosis.5,8,11–13 The natural history of pulmonary hypertension in infants with established chronic lung disease is unknown.

Pulmonary arterial pressure can be assessed non-invasively by Doppler echocardiography using pulmonary flow indices (and their calculated ratios) measured from the pulmonary artery Doppler velocity waveform. The ratio between the acceleration time (AT) and the right ventricular ejection time (RVET) is inversely related to PAP measured directly at cardiac catheterisation.14

The aim of our study was to use this technique to investigate changes in PAP during the 1st year of life in: (1) healthy preterm infants without respiratory disease, and (2) infants with established chronic lung disease.

Methods

Infants with chronic lung disease and healthy preterm infants were studied prospectively during a two year period between June 1995 and May 1997. Infants with chronic lung disease, defined as oxygen dependency for at least 28 days and beyond 36 post menstrual weeks in association with an abnormal chest radiograph, were studied at the regional chronic lung disease follow up clinic. Preterm infants without chronic lung disease, who had received mechanical ventilation for less than 24 hours and supplemental oxygen for less than 72 hours were studied as a healthy preterm reference population for comparison. Doppler echocardiographic assessments of PAP were performed at the time that the infants attended for routine follow up.

Pulmonary artery Doppler examination was performed using either a Hewlett Packard Sonos 100 or a Vingmed CFM 725 ultrasound system. A 7.5 MHz multifrequency imaging transducer combined with a 5 MHz Doppler transducer was used with each machine. After a preliminary cross sectional echocardiographic examination to exclude structural defects, a two dimensional image of the main pulmonary artery and pulmonary valve leaflets was obtained from a modified parasternal short axis...
### Table 1 Characteristics of study groups

<table>
<thead>
<tr>
<th></th>
<th>Infants with CLD (n = 54)</th>
<th>Healthy preterm infants (n = 44)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birthweight (g)*</td>
<td>1055 (510–2310)</td>
<td>1420 (840–2150)</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Gestation (weeks)*</td>
<td>28 (24–34)</td>
<td>30 (26–34)</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Sex (M:F)</td>
<td>28:26</td>
<td>23:21</td>
<td>1.0</td>
</tr>
<tr>
<td>Multiple pregnancy (%)</td>
<td>10 (19)</td>
<td>20 (45)</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Antenatal steroids (%)</td>
<td>45 (83)</td>
<td>31 (70)</td>
<td>0.25</td>
</tr>
<tr>
<td>Caesarean section (%)</td>
<td>32 (59)</td>
<td>27 (61)</td>
<td>1.0</td>
</tr>
<tr>
<td>Apgar score at 5 min*</td>
<td>8 (3–10)</td>
<td>9 (8–10)</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>Air leak (%)</td>
<td>11 (20)</td>
<td>11 (20)</td>
<td></td>
</tr>
<tr>
<td>Symptomatic PDA (%)</td>
<td>5 (9)</td>
<td>5 (9)</td>
<td></td>
</tr>
<tr>
<td>Total duration of ventilation (days)*</td>
<td>16 (1–64)</td>
<td>16 (1–64)</td>
<td></td>
</tr>
<tr>
<td>Duration of supplemental oxygen (weeks)*</td>
<td>39 (10–103)</td>
<td>39 (10–103)</td>
<td></td>
</tr>
</tbody>
</table>

*Values expressed as median (range).

CLD, chronic lung disease; PDA, patent ductus arteriosus.

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view. Doppler ultrasound examination was performed as described previously. Briefly, a Doppler signal was recorded from the centre of the main pulmonary artery immediately distal to the valve leaflets. Measurements of AT and RVET were made from the Doppler velocity waveform. Values of each were obtained from a minimum of five consecutive waveforms and the mean AT/RVET ratio calculated. Because there was no significant correlation between mean AT/RVET and R–R interval from the electrocardiograph, calculation of the corrected AT/RVET ratio was considered unnecessary. Subjective assessment of right ventricular contractility was normal in all patients. Doppler examinations were performed by a single observer (NVS) and infants were studied while asleep or during a period of quiet wakefulness. The intraobserver reproducibility of this technique has been reported previously.

Clinical and demographic data were collected from case notes. All infants who required mechanical ventilation for respiratory distress were treated with two doses of surfactant, immediately after birth and again at 12 hours of age. Symptomatic patent ductus arteriosus was recorded when an infant with a clinical diagnosis of patent ductus arteriosus was treated with indomethacin, or underwent surgical ligation. The total number of days of mechanical ventilation while on the neonatal unit, and the total duration of supplemental oxygen treatment were also recorded.

Supplemental oxygen treatment was prescribed to maintain arterial oxygen saturation (SpO2) consistently above 93%, measured using an Ohmeda Biox 3700e pulse oximeter. Oxygen was discontinued when an infant was able to maintain SpO2 values consistently above 93% during an overnight saturation study. Satisfactory SpO2 values (> 93%) were ensured throughout the Doppler assessment. All infants had stable oxygen requirements and no intercurrent respiratory illness at the time of study. Our study was approved by the local paediatric ethics committee and informed parental consent was obtained.

All statistical analyses were performed using SPSS for Windows, Release 6.0. Results are presented as median (range). Comparisons between infants with chronic lung disease and healthy preterm infants were carried out using the Mann-Whitney U test for continuous data, and χ² or Fisher’s exact test for categorical data. Serial measurements of AT/RVET were grouped and analysed at the following time periods: 36 weeks post menstrual age; at term; 1–13 weeks, 14–26 weeks, 27–39 weeks, and 40–52 weeks of age corrected for prematurity.

Correlation between changes in AT/RVET and age within individual infants (with and without chronic lung disease) was calculated using analysis of covariance. The overall rate of change in AT/RVET with time (AT/RVET velocity) was calculated for each infant using the first and last measurements made by using the following formula: change in AT/RVET divided by change in time, multiplied by 100 (and expressed as units/week). Median AT/RVET velocity was compared in healthy infants and infants with chronic lung disease.

Multiple linear regression analysis was used to determine which factors were independently associated with AT/RVET. Because multiple Doppler examinations had been performed two approaches were used:

1. Using all the available measurements, with AT/RVET as the dependent variable, the influence of birthweight, gestational age, sex, presence of chronic lung disease, and corrected age as explanatory variables was investigated.

2. In infants with chronic lung disease, with median AT/RVET as the dependent variable, the influence of birthweight, gestational age, duration of ventilation, duration of supplemental oxygen treatment, and median corrected age for each individual was investigated. The median values of AT/RVET and corrected age were used as summary measures of each variable in the regression model, to allow for multiple measurements in an individual.

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### Results

Fifty four infants with chronic lung disease and 44 healthy preterm infants were studied. Table 1 shows the demographic and clinical characteristics of the two groups. Infants with chronic lung disease were significantly lighter and less mature at birth, and had a lower 5 minute Apgar score. A significantly higher proportion of the healthy infants studied were born after a multiple pregnancy.

A total of 248 Doppler examinations were performed: 150 studies in infants with chronic lung disease and 98 studies in healthy infants. Table 2 summarises the timing and results of the examinations performed in each group. The median number of examinations was 2 (range, 2–6) in infants with chronic lung disease and 2 (range, 2–4) in healthy preterm infants. There was no significant difference between the two groups in the overall median corrected age at which the examinations were performed: 12 (range, 38 weeks post menstrual age to 44 weeks) in infants with chronic lung disease versus 13 (range, 38 weeks post menstrual age to 39 weeks) in healthy infants (p = 0.77).

The overall median AT/RVET was significantly lower in infants with chronic lung disease (table 2). Figure 1 shows changes in AT/RVET with time. Median AT/RVET in infants with chronic lung disease tended to be lower at all time points.
lower than that found in healthy infants throughout the period of study. This difference was significant at all time periods except at 36 weeks postmenstrual age and between 40 and 52 weeks of age corrected for prematurity. Within individual healthy preterm infants, changes in AT/RVET were significantly correlated with changes in time ($r = 0.67$; $p < 0.001$). There was a similar degree of correlation between the same variables in infants with chronic lung disease ($r = 0.55$; $p < 0.001$). There was also no significant difference in median AT/RVET velocity between the two groups (table 2).

In the first multiple regression model, AT/RVET was significantly and independently associated with the presence of chronic lung disease, gestational age, and corrected age at examination (table 3). In the second multiple regression model, AT/RVET was significantly independently associated with age at examination and inversely with duration of supplemental oxygen treatment in infants with chronic lung disease. The association with gestational age was of borderline significance (table 3).

**Discussion**

Pulmonary hypertension is common in preterm infants with respiratory distress syndrome and PAP remains raised in those who subsequently develop chronic lung disease.9 10 We evaluated changes in PAP, as assessed by the pulmonary artery Doppler derived AT/RVET ratio, in infants with established chronic lung disease and healthy preterm infants after discharge from hospital during the 1st year of life. Pulmonary arterial pressure is inversely related to age in both healthy infants and those with chronic lung disease. However, although PAP falls at a similar rate during infancy in both groups, infants with established chronic lung disease have persistently raised PAP compared with healthy preterm infants.

In healthy term infants, intra-acinar arterial multiplication parallels alveolar proliferation, which occurs most rapidly in the first 2 years of life. Intra-acinar vessels also increase in size with growth during infancy.13 This increase in the total cross sectional area of the pulmonary vascular bed results in an overall reduction in pulmonary vascular resistance. In childhood, this allows PAP to remain unchanged despite an increase in cardiac output.13

Lung development in non-ventilated preterm infants is appropriate for post menstrual age, suggesting that preterm birth is not itself associated with poor postnatal lung growth.20 In contrast, impaired alveolar and arterial development has been demonstrated in studies of preterm infants dying after mechanical ventilation in the early neonatal period.19 In our study, AT/RVET was significantly directly related to gestational age, indicating that PAP is higher in more immature infants in the 1st year of life, independently of the presence of chronic lung disease. Histological studies of the pulmonary circulation in infants and older children dying after the development of chronic lung disease have shown pronounced structural abnormalities, including increased medial thickness of small pulmonary arteries and distal extension of arterial smooth muscle.23 3 4 There is also a reduction in the number of peripheral arteries associated with a normal alveolar to arterial ratio, indicating failure of alveolar development.4 Disruption in growth and development of the pulmonary vascular bed might result in raised pulmonary vascular resistance and PAP in ventilated preterm infants who subsequently develop chronic lung disease.

Several cross sectional echocardiographic studies have described raised PAP in infants with chronic lung disease.21–23 Pulmonary hypertension with raised pulmonary vascular resistance has also been confirmed by direct measurement at cardiac catheterisation.4 12 13 However, the natural history of abnormalities of pulmonary haemodynamics in infants with chronic lung disease is not known. Reports in a few infants who have undergone repeat catheterisation have demonstrated a reduction in PAP with time, although PAP might remain persistently raised into later childhood.4 24 One infant, not receiving supplemental oxygen despite raised PAP, showed progressive pulmo-
nary hypertension in the 20 month period between catheter investigations. To our knowledge, there are no published studies of changes in PAP in healthy preterm infants or those with chronic lung disease during infancy after discharge from hospital.

Pulmonary arterial pressure and pulmonary vascular resistance in infants with chronic lung disease have been assessed non-invasively using M mode and Doppler echocardiography. The two methods most commonly used have been: (1) measurement of pulmonary flow indices, or (2) measurement of the maximal velocity of tricuspid regurgitation and application of the modified Bernoulli equation. Measurement of maximal tricuspid regurgitation velocity is generally accepted as the method of choice when assessing PAP non-invasively. However, the only study that has used this technique in infants with chronic lung disease found that the incidence of measurable tricuspid regurgitation was only 44%, despite sedation. The same study also reported a close correlation between measurements of AT/RVET and tricuspid regurgitation derived PAP. Pulmonary flow indices were chosen in our study because measurements could be performed successfully in almost all infants without the need for sedation.

Unlike previous studies, our study has compared PAP in infants with chronic lung disease with healthy preterm infants, rather than using a reference population of preterm infants who had received respiratory support but not developed chronic lung disease. This inevitably led to demographic differences between chronic lung disease infants and our reference group. Although it might have been preferable to have arranged Doppler assessments at predefined times, this was impractical in view of the geographical distribution of the study populations within the Mersey region. Instead, all assessments in our study were made on an opportunistic basis, at the same time as an infant attended follow up clinic. Infants with chronic lung disease who continued to attend clinic throughout the study period are likely to have been the sickest infants. This might have influenced our results through over representation of infants with low AT/RVET values towards the end of our study.

Reactivity of the pulmonary vascular bed to oxygen in infants with chronic lung disease has been confirmed in direct and indirect studies. However, the precise relation between different values of \( \text{SpO}_2 \) and PAP has not been fully evaluated. This might reflect the difficulty in maintaining a steady \( \text{SpO}_2 \) value during an echocardiographic examination. Because of the inherent inaccuracy in defining true \( \text{SpO}_2 \) on a background of continuous variation with time and movement artefact, all infants in our study were maintained at \( \text{SpO}_2 \) values above 93%, rather than attempting to define a precise \( \text{SpO}_2 \) value. It is possible that in some infants, maintaining an even higher \( \text{SpO}_2 \) value might have led to a reduction in pulmonary vascular resistance and PAP.

We have shown previously that persistently raised PAP at 24 hours of age is an important early predictor for the development of chronic lung disease in ventilated preterm infants. Our study has demonstrated an indirect independent relation between duration of supplemental oxygen treatment and AT/RVET. This suggests that PAP remains higher in infants who receive a longer period of oxygen treatment, and therefore presumably have more severe disease. Because none of our infants died within the period of our study, we were unable to examine the relation between PAP and mortality in our study group.

We have shown that although PAP is higher in infants with chronic lung disease than in healthy preterm infants, it tends to fall with increasing age during the 1st year of life at a similar rate in both groups. Pulmonary hypertension is also independently associated with a longer duration of supplemental oxygen treatment. Future research will need to consider the importance of pulmonary hypertension and the effects of pulmonary vasodilator treatment in influencing respiratory outcome in preterm infants with chronic lung disease.

NVS was supported by the British Heart Foundation (RF Martin Junior Research Fellowship).


