Decreased neutrophil apoptosis in tracheal fluids of preterm infants at risk of chronic lung disease

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Objective: To investigate the hypothesis that preterm infants who are more susceptible to lung damage have decreased neutrophil apoptosis, and to explore its relation to interleukin 10 (IL10) concentration.

Design: Prospective cohort design.

Patients: One hundred tracheal fluid specimens from 50 week-1 ventilated infants were examined for IL10 (by enzyme linked immunosorbent assay) and neutrophil apoptosis (by light microscopy).

Results: Neutrophil apoptosis was absent or less than 0.22% (median 0%) in the 11 infants with chronic lung disease (CLD) (24–31 weeks gestation) during the first 4 days of life. This was significantly lower than that of the 20 preterm infants without CLD (27–31 weeks gestation; median 0.47%, range 0–1.25%) and 19 term infants (median 0.5%, range 0–2.25%). There was an increase in apoptosis in infants with CLD (median 0.44%, p = 0.046) during days 5–7. Few infants without CLD were intubated beyond 4 days. Median apoptosis on days 5–7 was 0.26% and 2.78% for non-CLD preterm and term infants, but differences were not significant. IL10 concentration in tracheal fluid of infants with CLD was less than 5 pg/ml. None of the infants with IL10 greater than 5 pg/ml developed CLD. The range of IL10 concentrations in tracheal fluid from infants without CLD was wide (0–938 pg/ml). There was no apparent correlation between IL10 levels and percentage neutrophil apoptosis in infants without CLD.

Conclusion: Preterm infants with low levels of IL10 and neutrophil apoptosis may be predisposed to disordered lung repair. Further studies into the method of disposal of senescent neutrophils within preterm lungs are required.
a haemocytometer. Supernatant was stored at −80°C and examined in batches for IL8 and IL10. Two or more slides were made by cytopsin preparation of each sample, and the slides were stained with Diff-Quik (Lab Aids Pty Ltd, Narrabeen, Australia) and confirmed by agarose gel electrophoresis. This procedure involves the fixation of the cell pallet obtained immediately after centrifugation of the lavage material by 70% alcohol. To obtain DNA, the fixed cells were then treated by lysis and nuclear buffers. Ethidium bromide staining was used to ascertain if ladder development had occurred during agarose gel electrophoresis of the extracted DNA (concentration of agarose gel was 2%; electrophoresis was performed for 30 minutes using a Gene Power Supply (Pharmacia GPS 200/400) at 100 V).

Differential cell counts and characteristic changes in neutrophil apoptosis (chromatin condensation and simplification of nuclear structure) were assessed by phase contrast microscopy of 400 cells under oil immersion (fig 1). Even though the TUNEL (terminal deoxynucleotidyl transferase mediated dUTP nick end labeling) method is widely used to identify apoptotic cell fractions in cultured cell preparations, we used Diff-Quik staining to avoid some of the potential artifacts associated with the TUNEL assay—for example, dependence on staining kinetics of the reagent, or on tissue fixation, or on the extent of proteolysis of the cells.15

**Cytokine analysis**

IL10 was analysed in batches by enzyme linked immunosorbent assay (ELISA; R & D, Minneapolis, Minnesota, USA). The minimum detectable level of this assay was 0.5 pg/mL. The reproducibility of IL10 assays was confirmed by interlaboratory examination of seven selected tracheal fluid specimens (Airways Research Centre, John Hunter Hospital, Newcastle, Australia), as well as by the addition of anti-IL10 immunoglobulin and exogenous human IL10 to several selected specimens. Cytokine values were expressed as weight/volume of lavage because no other satisfactory correction factor for tracheal fluid cytokine measurement is currently available.14

**Statistical analysis**

This was performed with the Mann-Whitney U, Kruskal-Wallis, Student’s t, and Fisher exact tests. p < 0.05 was considered significant.

**RESULTS**

Of the 52 ventilated infants enrolled into our study, two died from necrotising enterocolitis before 36 weeks corrected age, and their results were excluded from our final analysis. Thirty one of the remaining 50 infants were preterm—that is, < 37 weeks gestation. Most (23/31, 74%) were ventilated for HMD. Table 1 summarises the clinical characteristics of all 50 infants. All HMD infants received at least one dose of exogenous bovine surfactant. None of the infants received corticosteroids during the first week even though most infants with CLD (8/11, 73%) and a term girl with MAS were treated with dexamethasone during or after the second week of life.

Two mothers had chorioamnionitis and both their infants developed CLD. Placental bacterial cultures were sterile in one and positive for group B streptococcus in the other. In our institution, low vaginal swabs are routinely performed on gravid women either at 28 weeks gestation or at delivery (if birth occurred before 28 weeks). Group B streptococcus was grown in the low vaginal swabs of four women (three with infants with CLD, and one with a term infant), and these women were treated with intrapartum penicillin. None of their infants developed positive blood cultures. A non-CLD preterm infant grew Enterococcus colit on blood collected immediately after birth. Her mother's low vaginal swab was negative.

The 11 (36%) preterm infants who developed CLD were significantly more premature than the 20 preterm infants whose acute lung disease resolved, and the duration of ventilation was significantly longer (table 1). Two 28 week preterm infants without CLD were re-intubated at 52 and 21 days of age because of bronchiolitis and coagulase negative staphylococcus sepsis respectively. These infants were not oxygen dependent and both were easily extubated (into room air) after resolution of the acute illnesses.

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p = 0.046) or the combination of all preterm infants (median 0.26%, range 0–1.5%; p = 0.019). The highest (10.5%) was from a female infant with MAS from tracheal fluid specimens obtained on day 5. Figure 2 summarises the results.

As most of the infants without CLD were extubated after the first 4 days of life, only a few tracheal fluid specimens were obtained from these infants during this period. Tracheal fluid results were thus divided into days 1–4 and 5–7 for comparison. Some infants had two tracheal fluid specimens in either of the time epochs. To achieve equal weight for each infant in this analysis, the mean value of these was taken as the single individual value within the time epoch. Results remained consistent with the trend illustrated in fig 2. Neutrophil apoptosis was either absent or less than 0.22% in the 11 infants with CLD during the first 4 days of life (median 0%). Compared with infants without CLD during the first 4 days, those with CLD had a lower percentage of apoptotic neutrophils (non-CLD preterm: median 0.47%, range 0–1.25%, p = 0.001, Mann-Whitney; term infants: median 0.5%, range 0–3.25%, p = 0.018). There was a trend towards an increase in the percentage of neutrophil apoptosis in infants with CLD during days 5–7 (median 0.44%, range 0–1.3%, p = 0.046). Median apoptosis on days 5–7 for non-CLD preterm and term infants was 0.26% and 2.78% respectively, but differences did not reach significance.

Neutrophil apoptosis in relation to IL10

IL10 concentration was less than 5 pg/ml in all CLD tracheal fluid specimens (fig 3). The range of IL10 from the non-CLD preterm specimens was much wider (0–938 pg/ml) and fluctuated throughout the first week of life. IL10 levels were greater than 5 pg/ml in 11 non-CLD preterm infants and in six term infants. Respiratory conditions in this group of infants were varied and ranged from those necessitating profound ventilatory support (such as term infants with MAS) to those with only mild disease (such as transient tachypnoea of the newborn) or no lung disease (such as infants intubated electively for seizure control).

The relation between IL10 and apoptosis was examined in non-CLD preterm and term infants because IL10 and neutrophil apoptosis levels were both low in infants with CLD. There was no apparent correlation between IL10 and apoptosis in either preterm (r = 0.128, p = 0.357, NS) or term (r = 0.212, p = 0.331, ns) infants.

DISCUSSION

To our knowledge, our study is the first to examine both reduced neutrophil apoptosis and IL10 in the tracheal fluid of premature infants who subsequently develop CLD. IL10 inhibits the production of proinflammatory cytokines such as IL8, tumour necrosis factor (TNF)α, and IL6 from neutrophils and macrophages. These cytokines are essential for maintaining inflammation. IL8, for example, is a powerful pulmonary neutrophil chemotaxin that acts in concert with TNFα to prolong neutrophil longevity by inhibiting apoptosis. In a recent pilot study, we found no difference in tracheal fluid IL8 between infants with and without CLD, although IL8 tended to decrease towards the end of the first week in infants without CLD. There is, as yet, little evidence that neutrophils entering the lung as a result of injury are able to return to the microcirculation after dissipation of the injurious process. Therefore, for healing to occur, redundant neutrophils need to be inactivated and cleared from the lung.

### Table 1 Basic and clinical data of study infants

<table>
<thead>
<tr>
<th></th>
<th>CLD (n=11)</th>
<th>Preterm non-CLD (n=20)</th>
<th>Term (n=19)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age (weeks)*</td>
<td>27 (24–31)</td>
<td>31 (27–33)</td>
<td>40 (38–42)</td>
</tr>
<tr>
<td>Birth weight (g)*</td>
<td>813 (520–1500)</td>
<td>1600 (830–2190)</td>
<td>3640 (2220–4440)</td>
</tr>
<tr>
<td>Male</td>
<td>9 (82%)</td>
<td>14 (70%)</td>
<td>11/19 (58%)</td>
</tr>
<tr>
<td>Days ventilated*</td>
<td>30 (4–90)</td>
<td>5.5 (2–10)</td>
<td>5 (2–11)</td>
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<tr>
<td>Indications for ventilation</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>HMD</td>
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<td>14</td>
<td>5</td>
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<tr>
<td>TTN</td>
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<td>4</td>
<td>2</td>
</tr>
<tr>
<td>Apnoea</td>
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<td>2</td>
<td></td>
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</tr>
<tr>
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</tr>
<tr>
<td>CHD</td>
<td>0</td>
<td>0</td>
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</tr>
<tr>
<td>Sepsis</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
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</table>

Values are median (range) or number (%).

* p<0.001, Kruskal-Wallis.

CLD, Chronic lung disease; HMD, hyaline membrane disease; TTN, transient tachypnoea of the newborn; PPHN, persistent pulmonary hypertension; CHD, congenital heart disease; HIE, hypoxic ischaemic encephalopathy.
Apoptotic neutrophils lose the ability to degranulate potentially injurious intracellular contents.30 Macrophages that phagocytose apoptotic neutrophils do not release proinflammatory agents such as IL8, IL1β, and TNFα.21–23 Neutrophil apoptosis is thus an important mechanism by which various forms of acute lung injury are resolved.31–33 In this study, we hypothesised that premature infants were more susceptible to disordered lung healing because of reduced neutrophil apoptosis. Grigg et al.34 noted apoptotic neutrophils at single time points in the lungs of eight mechanically ventilated infants. Their results, however, did not correlate with pulmonary outcome. We examined ventilated infants on multiple occasions over the first week of life and showed that the percentage of apoptotic neutrophils in tracheal fluid did not increase above 0.25% of the total neutrophil count until after the fourth day of life in infants with CLD. Our results suggest that this relative delay may expose vulnerable infants to continued neutrophil mediated lung injury.35 Periods of specimen analysis and collection were divided into two periods of the first week of life—that is, days 1–4 and days 5–7—because fewer tracheal fluid specimens were collected after the first 4 days of life in non-CLD preterm and term infants and because analysis of neutrophil counts in the later part of the week would have allowed us to capture the persistent pulmonary neutrophilia characteristic of developing CLD.25,36

IL10 is currently the only known endogenous factor that promotes neutrophil apoptosis.37 Exogenous IL10 attenuates pulmonary inflammation in experimental animals by accelerating neutrophil apoptosis and decreasing neutrophil influx.25 Various forms of chronic and severe pulmonary inflammatory disorders are associated with low levels of IL10, including cystic fibrosis, asthma, and fatal acute respiratory distress syndromes.26–28 IL10 levels are also reduced early in life in infants who develop CLD of prematurity.26–28 McColm et al.39 studied 11 infants less than 30 weeks gestation, nine of whom developed CLD. Detectable IL10 was found in 57% of the tracheal fluid specimens regardless of subsequent CLD development. However, only two infants without CLD were studied.

and higher gestation infants were not studied by their particular laboratory methods for comparison. We have shown (in a concurrent study) that, although IL10 is predominantly governed by gestation, more mature infants who were not expected to develop CLD also had minimal IL10 in their lung fluid. For example, a 29 week male infant needing only minimal ventilation for apnoea and a 31 week male infant with mild HMD had neither IL10 or apoptotic neutrophils in any of their week 1 tracheal fluid specimens. Nevertheless, higher levels of IL10 produced by more mature infants may have contributed to lung healing by constraining neutrophil activity, as evidenced by increased levels of apoptosis. This study, however, failed to show a positive correlation between IL10 and neutrophil apoptosis.

Other cytokines, however, also significantly influence neutrophil longevity, and it is difficult to determine from our study whether a causal relation exists between endogenous IL10 and neutrophil apoptosis. Proinflammatory cytokines such as IL8 and IL6 retard neutrophil apoptosis and are also increased in the lung fluid of CLD prone infants.37–50 There were also infants in our study who did not develop CLD despite having neither high levels of IL10 nor apoptotic neutrophils. Most of these infants, nevertheless, were more mature, and the pulmonary protective value of IL10 may be only significant in the most at risk—that is, the extremely preterm population.

The peak frequency of neutrophil apoptosis occurs 6–72 hours after pulmonary injury,31 and previous observations have shown that neutrophil apoptosis and subsequent phagocytosis by primed macrophages is a rapid process; neither neutrophils nor their cytoplasmic contents are recognisable as soon as 30 minutes after engulfment.31 Optimal in vitro identification of neutrophil apoptosis in our study may have been limited by various clinical constraints (such as laboratory availability and tracheal suction timing). Non-CLD specimens in our study were less complete than CLD specimens because most of these infants had been extubated by the end of the week. Therefore, we do not have a complete profile of infants without CLD throughout the first week, and IL10 and neutrophil apoptosis may have been evident in these infants if sampling times had differed. Even so, our findings describe a novel lack of neutrophil apoptosis and IL10 in CLD specimens compared with non-CLD specimens during the first 4 days of life.

We identified neutrophil apoptosis by light microscopy in a similar fashion to Savill et al.37 by noting shrinkage of the entire neutrophil and condensation of the nuclear chromatin into a rounded lobe. We also observed some atypical clump-like changes suggested by Ishii et al.34 to be representative of an early part of the apoptotic process.

Exogenous factors such as commonly used medications may modulate the rate of neutrophil apoptosis. Erythromycin38 and theophylline39 accelerate, whereas glucocorticoids retard,39 neutrophil apoptosis. The anti-inflammatory properties of glucocorticoids are well known. Despite retarding neutrophil apoptosis, glucocorticoids promote macrophage phagocytosis of apoptotic white cells.40 However, none of the infants in our study were treated with glucocorticoids during week 1, and most of the infants without CLD (term and preterm) did not receive theophylline until immediately before extubation. We therefore cannot conclude that the difference in the frequency of apoptosis between the groups of infants in our study was influenced in any way by theophylline or glucocorticoids. Further study is needed to assess the effect that these medications have on early pulmonary inflammation in susceptible preterm infants.

In conclusion, our results show that neutrophil apoptosis increases with gestational maturity and that infants who subsequently developed CLD had negligible amounts of apoptosis and IL10. However, the interplay of factors controlling pulmonary inflammation is complex and probably not dominated by any one cytokine. IL10 has anti-inflammatory properties that...
Neutrophil apoptosis in chronic lung disease

may have the potential to mitigate the neutrophil mediated disorganised pulmonary healing and repair characteristic of CLD. Our study supports the observation of Jones et al27 showing low IL10 in tracheal fluid in CLD prone infants. As glucocorticoids, with all their numerous side effects, are the only current relatively beneficial treatment for CLD, further investigations into ways of modulating the inflammatory profile of the preterm lung with novel anti-inflammatory treatments during the first week of life may eventually help to prevent and treat CLD.

ACKNOWLEDGEMENTS
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REFERENCES