NF-κB in tracheal lavage fluid from intubated premature infants: association with inflammation, oxygen, and outcome

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**Objectives:** To determine if tracheal lavage concentrations of the transcription factor NF-κB, which is activated by risk factors associated with bronchopulmonary dysplasia (BPD) and induces expression of cytokines associated with BPD, is related to BPD in premature infants.

**Design:** Serial tracheal lavage samples from intubated premature infants were analysed for cell count and concentrations of interleukin (IL)8 and NF-κB, corrected for dilution by secretory component concentrations.

**Setting:** Level III university hospital neonatal intensive care unit.

**Patients:** Thirty three intubated infants (mean (SD) birth weight 903 (258) g, median gestation 27 weeks (range 24–31)) in the first 14 days of life.

**Main outcome measures:** Tracheal effluent NF-κB, IL8, and cell counts, corrected for dilution by secretory component measurement.

**Results:** Square root transformed NF-κB concentrations were significantly related to signs of inflammation (cell count, p = 0.002; IL8, p = 0.019) and to simultaneous fraction of inspired oxygen in samples from the first 3 days of life (r = 0.512, p<0.003). Of the 32 subjects with samples in the first 3 days of life, the half who either died or had BPD had higher NF-κB concentrations than those without BPD (square root concentration 0.097 (0.043) v 0.062 (0.036) μg/μg protein/μg secretory component, p = 0.018).

**Conclusions:** Tracheobronchial lavage NF-κB concentrations are related to lung inflammation, oxygen exposure, and pulmonary outcome in intubated preterm infants. NF-κB activation may be an early critical step leading to BPD.

Bronchopulmonary dysplasia (BPD) occurs primarily in premature infants who are mechanically ventilated and exposed to oxygen early in their life. It is associated with early inflammation of the airways and lung interstitium, mediated by proinflammatory cytokines (interleukin (IL)1β, IL6, IL8, monocyte chemoattractant protein (MCP)1, and intercellular adhesion molecule (ICAM)1), which are raised in tracheobronchial lavage obtained from patients who progressed to BPD. Expression of these mediators is partially controlled through the activation of transcription factors. These are proteins that, on appropriate stimulation, attach to a recognition sequence in the promoter region of specific genes. One of these transcription factors is nuclear factor-kappa B (NF-κB).

NF-κB, a ubiquitous transcription factor, promotes the expression of many genes, including proinflammatory cytokines associated with the development of BPD. It is activated by a variety of factors, including infectious stimuli, inflammatory cytokines, deformation, oxidants, and other causes of cell stress. It has roles in controlling apoptosis and cell proliferation and differentiation and may be involved in lung development.

Activation of NF-κB occurs in many inflammatory conditions both in vivo and in vitro. Lung macrophages from adults at risk of acute respiratory distress syndrome had significantly more activated NF-κB than from those who did not develop acute respiratory distress syndrome. NF-κB activation was higher in monocytes and neutrophils in adults with fatal systemic inflammatory response than in those who survived.

As NF-κB is activated by stimuli associated with the development of BPD, promotes the expression of proinflammatory mediators that are increased in the lungs of infants who develop BPD, and has been found in older patients with acute respiratory distress syndrome, we hypothesised that activation of NF-κB may be a critical early step in the pathophysiology of BPD. The two aims of this study were to see if there is increased activation of NF-κB in tracheal lavage effluent cells from premature infants who develop BPD and to examine if clinical conditions that might activate NF-κB are related to potential inflammatory consequences of its activation.

**METHODS**

**Subjects and sampling**

Subjects who were <1500 g birth weight and intubated in the first 3 days of life were eligible for enrolment. Exclusion criteria were likely demise and lack of informed consent. Prophylactic surfactant therapy was given to inborn neonates <30 weeks gestation, and rescue surfactant provided to all others who were diagnosed with respiratory distress syndrome. Infants weighing <1000 g at birth were usually placed on high frequency oscillatory ventilation. Prophylactic indomethacin was given to babies <30 weeks gestational age. The study was approved by the institutional review board, and informed consent was obtained from parents before any sampling.

Tracheal lavage fluid during routine suctioning was collected on days 1–2, 3, 5, 7, and 14 of life if the patient was still intubated. Material was collected using previously

**Abbreviations:** BPD, bronchopulmonary dysplasia; FiO2, fraction of inspired oxygen; IL, interleukin; NF-κB, nuclear factor-kappa B; SC, secretory component.
published methods. The samples were transported to the laboratory on ice and processed immediately.

A manual white blood cell count was performed. After centrifugation (1200 g for five minutes at 4°C), protease inhibitors (1 M 0.5 M dithiothreitol, 5 M 100 mM phenylmethanesulphonyl fluoride, 20 M 1 mg/ml leupeptin, 10 M 5 mg/ml aprotinin, 10 M 100 mM benzamidine; all from Sigma, St Louis, Missouri, USA) were added per ml supernatant, and the pelleted cells and supernatant stored at −80°C.

Assays

Nuclear protein was isolated from the thawed cell pellet in the three step process according to the manufacturer’s instructions (Nuclear Extract Kit; Activemotif, Carlsbad, California, USA). The protein concentration was determined by Coomassie protein assay. NF-kB in nuclear protein extract was determined by enzyme linked immunosorbent assay (TransAM NF-kB p65; Activemotif), again following the manufacturer’s instructions. Assay specificity was confirmed using the provided competitive oligonucleotides. The provided HeLa cell extract was used to produce a standard curve.

Equal amounts of nuclear protein were used, and the results normalised to both the amount of nuclear protein and the amount of secretory component (SC) and expressed as ng/μg nuclear protein/μg SC.

The supernatant was assayed for cytokine IL8 by enzyme linked immunosorbent assay (IL8 DuoSet; R & D Systems, Minneapolis, Minnesota, USA), diluted at a 1:1 to 5:1 diluent to sample ratio using standard diluent to bring the sample concentration into the assay range 25–2000 pg/ml. Samples were run in duplicate. IL8 concentration was expressed as pg/μg SC.

The concentration of SC in the specimen supernatant was determined using a quantitative sandwich enzyme linked immunosorbent assay. The capture antibody (mouse monoclonal anti-hSC; NI 194-4), detection antibody (horseradish peroxidase conjugated polyclonal goat anti-hSC), and standard (free SC from human milk) were purchased from Accurate Chemical, Westbury, New York, USA.

Clinical data and analysis

Data on perinatal factors, including infection and steroid use, and concurrent ventilator settings were collected prospectively. The fraction of inspired oxygen (FIO2) and ventilatory settings at the time of sampling were used. BPD was defined as a need for supplemental oxygen at 36 weeks postconceptional age (BPD group). The median amount of NF-kB in the tracheal lavage samples was 4.51 ng/μg protein/μg SC (5th–95th centile range 1.25–27.7). Cell count and IL8 concentration were 1.35 cells/μg SC (5th–95th centile range 0.11–6.32) and 4.54 pg/μg SC (5th–95th centile range 0.13–19.29) respectively.

There was no significant difference between the NF-kB concentrations in the first 3 days of life from the two subjects whose mothers did not receive prenatal steroids compared with the subjects whose mothers were treated (square root 0.043 (0.007) v −0.067 (0.045) μg/μg protein/μg SC, p = 0.425). Similarly, samples from the three subjects with culture proven sepsis during the sampling period did not differ significantly from those obtained on the same day from subjects without infection (square root infected 1.47 (0.59) v non-infected 1.60 (0.42) μg/μg protein/μg SC, p = 0.733). Postnatal steroids were given to five subjects during the study period, one on days 3–5, one on days 6–10, and three on days 11–13. The last three subjects had mean square root transformed NF-kB lavage concentrations on day 14 of 1.31 (0.61) compared with 0.98 (0.43) μg/μg protein/μg SC in the three subjects who had not received steroids (p = 0.487). It must be noted that, owing to the relatively small number of samples in each condition, the possibility of making either a type I or type II error regarding the effect of infection or steroids is significant.

There were significant relations between the sample NF-kB concentration and concurrent inflammatory marker (cell count, F = 11.56 p = 0.002) and mediator (IL8, F = 6.06, p = 0.0194) in all available lavage samples. The relation between NF-kB and one potential stimulant, oxidant stress, was evaluated by analysis of NF-kB concentration in a lavage sample with oxygen concentration at the time of sampling, but was not significantly related using the mixed effects model analysis (FIO2, F = 0.82, p = 0.372).

To examine whether early NF-kB activation precedes development of BPD, we focused on the samples obtained in the first 3 days of life and compared the values from subjects who survived without BPD with those from subjects who either died or developed BPD. If a subject had more than one sample obtained during this early period, the results were averaged. In this early set of samples, the transformed NF-kB concentration correlated with FIO2 (r = 0.484, p = 0.005), but did not reach the stricter threshold for statistical significance—0.391, p = 0.033 or IL8 (r = 0.375, p = 0.035). There was a significant correlation between the individual transformed NF-kB concentrations from days 1–3 and the transformed IL8 concentration at the subsequent sample—that is, day 1 or 3 NF-kB and day 3 or 5 IL8 (n = 14, r = 0.572, p = 0.033).

Of the 32 subjects, two died and 14 were oxygen dependent at 36 weeks postconceptional age (BPD group). The

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remaining 16 were classified as no BPD. Table 1 compares these two groups. As expected, those in the BPD group were smaller and younger. The groups did not differ with respect to lavage cell count or IL8 concentration. In contrast, NF-κB concentration was higher in the BPD group (0.097 (0.043) vs. 0.062 (0.036) μg/g protein/μg SC, p = 0.018). However, when gestational age was taken into account using multiple logistic regression, NF-κB activation was no longer significant.

**DISCUSSION**

Inflammation, seen early in BPD, is a complex response to external stimuli. The transcription factor, NF-κB, plays a major role in transducing the stimulus signal(s) into the mediators that promote, modulate, and inhibit the inflammatory response. When activated by environmental stimuli such as shear stress,14 oxidants,15 or particles,16 or through ligands such as endotoxin, IL1β, or tumour necrosis factor α attaching to cell surface receptors,17 the NF-κB dimer is translocated to the nucleus where it attaches to a specific DNA sequence in the promoter region to help induce gene expression. Many of the gene products associated with NF-κB have also been associated with BPD.18–20 In addition, many risk factors for BPD (volutrauma,20 infection,20 and oxygen21) can activate NF-κB. In clinical series, NF-κB activation has been found in lung cells of patients with acute respiratory distress syndrome,19 chronic obstructive pulmonary disease,22 lung transplant,23 and asthma,24 and the degree of NF-κB activation correlates with disease severity.

In this study, we examined two questions. Firstly, we found that there was a significantly higher NF-κB concentration in patients who died or developed BPD than in those who did not. That this difference was influenced by gestational age is consistent with the fact that prematurity is the strongest risk factor for BPD and that inflammation is one of the earliest processes found in those who develop BPD. Secondly, we looked at the relation between the NF-κB concentration in the lavage fluid and other factors in the sample, and between the NF-κB concentration and conditions at the time of sampling to examine clinically some of the possible causes and effects of NF-κB activation. NF-κB concentration was related to the amount of oxygen exposure at the time of sampling, consistent with the fact that oxidant stress activates NF-κB. The relation between the transcription factor and an inflammatory mediator, IL8, also agrees with what is known about the consequences of NF-κB activation.

The identity of the cells from which the NF-κB was extracted was not established in this study because diverting even a small number of cells for a differential cell count might have reduced the yield of nuclear protein below the assay detection limit. As we do not know if these cells were resident—for example, macrophages—or migrated to the lungs—for example, neutrophils—these results, while consistent with the hypothesis that NF-κB activation leads to lung inflammation and BPD, cannot prove it.

If confirmed, our observations point towards an earlier step in the processes that translate noxious stimuli into inflammation and, ultimately, BPD in the youngest, most vulnerable patients. To determine that NF-κB activation is, in fact, a critical component of this pathway will require studies in

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Clinical characteristics and lavage variables on days 1–3 for the group of infants who did not develop bronchopulmonary dysplasia (No BPD) and the group who died or did develop BPD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Variable</td>
<td>No BPD</td>
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<td>Total</td>
<td>16</td>
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<tr>
<td>Died</td>
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<tr>
<td>Birth weight (g)</td>
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<tr>
<td>Gestation (weeks)</td>
<td>28 (25.3–30.7)</td>
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<td>Days in O2</td>
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<td>Lavage IL8 (μg/μg SC)</td>
<td>2.41 (1.68)</td>
</tr>
<tr>
<td>Lavage cell count (cells/μg SC)</td>
<td>1.125 (0.735)</td>
</tr>
<tr>
<td>Lavage NF-κB (pg/μg protein/μg SC)</td>
<td>0.062 (0.036)</td>
</tr>
</tbody>
</table>

Values are mean (SD) or median (5th–95th centile). The lavage variables are square root transformed.

*Does not include the two subjects who died.

IL, interleukin; NF-κB, nuclear factor-kappa B; SC, secretory component.
which inflammation and BPD are prevented when NF-κB activation is blocked. Several studies have provided some preliminary data supporting this in animals and in vitro.

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