Association of fever and severe clinical course in bronchiolitis

A Sahib El-Radhi, William Barry, Swatee Patel

Abstract
Little attention has been given to the relation between fever and the severity of bronchiolitis. Therefore, the relation between fever and the clinical course of 90 infants (59 boys, 31 girls) hospitalised during one season with bronchiolitis was studied prospectively. Fever (defined as a single recording > 38.0°C or two successive recordings > 37.8°C) was present in 28 infants. These infants were older (mean age, 5.3 ± 4.0 months), had a longer mean hospital stay (4.2 ± 2.7 days), and a more severe clinical course (71.0% ± 29.0%) than those infants without fever. Radiological abnormalities (collapse/consolidation) were found in 60.7% of the febrile group compared with 14.8% of the afebrile infants. These results suggest that monitoring of body temperature is important in bronchiolitis and that fever is likely to be associated with a more severe clinical course and radiological abnormalities.

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Keywords: fever; bronchiolitis; disease severity

Bronchiolitis is a major cause of lower respiratory tract infection in infancy. Most children have had respiratory syncytial virus (RSV) infection by the age of 2 years.1 2 Pre-existing chronic lung disease, congenital heart disease, immunodeficiency, prematurity, parental smoking, and infection in very early infancy predispose to severe illness and occasional death.3–5 Apnoea or respiratory arrest before hospitalisation and pulmonary consolidation on chest x ray have been associated with increased morbidity.6 A greater disease severity has been reported in group A RSV than group B infection.7

There are few reports of the incidence of fever in bronchiolitis, or its relation to the severity of the disease or radiological findings; furthermore, any such reports have produced conflicting results. Fever has been variously described as being very common in the initial phase of the illness, but largely disappearing by the time of hospitalisation8; as a characteristic feature of the disease9; as intermittent and rarely exceeding 38–39°C10; or as a sign rarely exceeding 38°C.11 These studies have merely stated the incidence of fever but we are not aware of the evidence underlying these conclusions. Those few studies that have investigated the incidence of fever in bronchiolitis have involved small numbers of children. A Swedish study12 of 15 infants with RSV infection reported fever in 14. Another study13 reported an incidence of fever as low as 12% among 25 infants aged less than 3 months with RSV infection. In addition, this study found that all six children with bronchiolitis caused by parainfluenza virus were afebrile.

Information on the relation between fever and clinical severity is even more limited. A single report14 on the clinical findings and severity of bronchiolitis showed that respiratory rate on admission, chest wall indrawing, temperature, heart rate, and liver displacement did not predict the severity of the infection. However, of the six clinical signs studied to predict oxygen requirements, fever was not included. More information is required to determine whether there is an association between the severity of bronchiolitis and fever.

Our study was carried out to assess the extent of fever in bronchiolitis, whether the clinical course differs between febrile and afebrile infants with bronchiolitis, and whether fever in bronchiolitis is beneficial or harmful.

Patients and methods
All infants aged over 1 month who were admitted to Queen Mary’s Hospital, Sidcup, UK during one season with a diagnosis of acute bronchiolitis were studied prospectively. Patients were admitted between November 1997 and February 1998. In our hospital, between 80 and 120 infants are admitted with the disease each winter. The diagnosis of bronchiolitis was made in the presence of a history of upper respiratory tract infection followed by acute onset of respiratory distress with cough, breathlessness, and wheeze, and clinical signs of chest overinflation, tachypnoea, rhonchi, or crepitations occurring during a winter epidemic of bronchiolitis.15 Only children with their first episode of bronchiolitis were included.

The timing of discharge was based on the establishment of a normal feeding pattern without an oxygen requirement and absence of fever, in accordance with our usual practice.

The severity of the infection was assessed using a simple severity index as described by McIntosh et al.16 This index takes the requirement of oxygen to be the best single initial measurement of the severity of the illness in bronchiolitis. Infants who required mechanical ventilation were classified as very severe, those who required oxygen supplement as severe, and those who required admission for observation without oxygen requirement as mild.

The axillary temperature was recorded on admission and monitored at least four hourly. Fever was defined as a single temperature recording > 38°C or two successive
temperature recordings > 37.8°C, taken four hours apart during the first 24 hours of admission. All children underwent examination of the nasopharyngeal aspirates for RSV and had a chest x ray. RSV was detected using Directigen RSV (Becton Dickinson, Oxford, UK) kits. Results were given as positive, negative, or uninterpretable and, in the latter case, a repeat measurement was carried out using an appropriate dilution. The RSV antigen was the only virus tested for.

A chest x ray was taken either on admission or within the first 24 hours of admission, and some x rays were repeated in cases of clinical deterioration. All x rays were reported upon by a radiologist for review after the discharge of the infants. The radiologist was unaware of the clinical severity or body temperature, but aware of the clinical diagnosis. The films were examined for the presence of hyperinflation and its degree, peribronchial streaky infiltration, and segmental/lobar collapse, with or without consolidation.

All children were managed according to a routine protocol, which included isolation in cubicles, continuous monitoring of oxygen saturation by pulse oximetry, and the provision of supplemental oxygen if required to maintain oxygen saturation > 92%. Breast milk or formula milk was offered if tolerated, with appropriate volume for age. Children not tolerating oral feeds because of dyspnoea, or those with severe bronchiolitis at presentation, received intravenous fluid and/or milk via a nasogastric tube. Nebulised ipratropium bromide was often prescribed as a trial and continued in those who responded. Antibiotics were administered if there was radiological evidence of collapse/consolidation and to those children who appeared toxic.

The t test (unrelated) and χ² test were used for statistical analysis.

**Results**

Ninety infants (59 boys, 31 girls) with a mean age of 4.4 months (SD, 2.7; range, 1–11.75) were studied. The duration of their illness from onset of coryzal symptoms until admission ranged between one and 13 days. The duration of their stay in hospital ranged from one to 10 days, with a mean of 3.1 days (SD, 2.7). A history of fever was present in 25 children and this information was obtained from the parents. RSV was identified from the nasopharyngeal aspirates in 55 of the 90 children tested.

The children were divided into two groups according to body temperature: febrile and afebrile. Table 1 shows the clinical data and results of radiological abnormalities. The mean age of the febrile group was significantly higher than that of the afebrile group (p = 0.033; 95% confidence intervals [CI], 0.1 to 2.5). The mean length of hospital stay was significantly higher in the febrile group (4.2 vs 2.7 days; p < 0.005; 95% CI, 0.7 to 2.3). A significantly higher disease severity was seen in the febrile group (71.0%) compared with the afebrile group (29.0%; p < 0.005). One child was classified as having very severe disease in the afebrile group. No deaths occurred. Thirteen infants were born prematurely, four in the febrile group and nine in the afebrile group. Similar proportional results were obtained from RSV testing in the febrile and afebrile groups. There was no known case with underlying cardiac or pulmonary disease or immunodeficiency.

There were 28 febrile children, with most body temperature measurements ranging between 38°C and 39°C (table 2). Eighteen children had a history of fever in association with an upper respiratory tract infection before admission. Children with a body temperature > 39°C (n = 17) tended to be older, had a similar mean duration of stay in hospital, and had similar clinical severity, although they were more likely to have radiological abnormalities compared with those with a body temperature < 39°C (p = 0.066). The mean duration of fever was 2.2 days (range, 0.5–5; SD, 1.3).

Overall, 20 of the 28 febrile children had a severe clinical score. The remaining 62 children were afebrile on admission and remained so throughout their hospital stay. A history of fever before admission was obtained from seven of these infants. Only one child had a very severe clinical course and required ventilation for two days. He made a full recovery.

Because the radiological changes of collapse and consolidation are virtually indistinguishable, both were grouped together (table 1). The incidence of segmental/lobar collapse/consolidation was significantly higher in the febrile group than in the afebrile group.

**Table 1** Clinical data of 90 children with bronchiolitis

<table>
<thead>
<tr>
<th>Groups</th>
<th>Mean age in months (range)</th>
<th>Mean length of stay in days (range)</th>
<th>Clinical severity</th>
<th>Radiological changes (segmental/lobar collapse/consolidation)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Severe (%)</td>
<td>Mild (%)</td>
</tr>
<tr>
<td>Febrile (n = 28)</td>
<td>5.3 (1–10.25)</td>
<td>4.2 (1–13)</td>
<td>20 (71.4)</td>
<td>8 (28.6)</td>
</tr>
<tr>
<td>Afebrile (n = 62)</td>
<td>4.0 (1–11.75)</td>
<td>2.7 (1–10)</td>
<td>18* (29)</td>
<td>44 (71)</td>
</tr>
<tr>
<td>p Value</td>
<td>0.033</td>
<td>&lt; 0.005</td>
<td>&lt; 0.005</td>
<td>&lt; 0.005</td>
</tr>
</tbody>
</table>

*One child had a very severe clinical course (ventilated) in the afebrile group.

**Table 2** Ranges of body temperature in 28 febrile children

<table>
<thead>
<tr>
<th>Body temperature (°C)</th>
<th>Children (n)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 38.0</td>
<td>1</td>
</tr>
<tr>
<td>&lt; 39.0</td>
<td>16</td>
</tr>
<tr>
<td>&lt; 40.0</td>
<td>9</td>
</tr>
<tr>
<td>≥ 40.0</td>
<td>2</td>
</tr>
<tr>
<td>Total</td>
<td>28</td>
</tr>
</tbody>
</table>
Fever and severe bronchiolitis

Discussion

The mean age of the population studied was 4.4 months, which is similar to that reported elsewhere, indicating that bronchiolitis mainly affects infants aged 2–5 months. The mean length of hospitalisation of 3.3 days is, however, significantly shorter than that reported from several hospitals in Canada (8.6–11.8 days), but is closer to that reported from Manchester, UK (4.5 days). Local variability influencing the pattern and severity of RSV infection, and thus the length of hospitalisation, has been reported.

One of the limitations of our study was virological testing for RSV only. It would have been interesting to investigate differences in the incidence of fever and disease severity among other viruses that cause bronchiolitis, such as influenza and parainfluenza viruses, adenoviruses, and rhinoviruses. As previously noted, one study found that all six children with bronchiolitis caused by parainfluenza virus were afebrile.

Dual infection with other viruses, such as adenoviruses, or organisms such as Chlamydia trachomatis or Mycoplasma pneumoniae, is common and has been found in at least 5–10% of cases of lower respiratory tract infection infected with RSV.

The main objective of our report was to study the clinical sign of fever in children with bronchiolitis. Fever is the most common sign of infectious diseases. Studies are needed to determine in specific paediatric diseases whether fever alone is beneficial, harmful, or without effect. In some diseases—for example, salmonella gastroenteritis, the presence of fever was associated with a shorter duration of convalescent faecal bacterial excretion than was seen in afebrile children. Children with high fever at the onset of the initial febrile convulsion had a significantly decreased recurrence rate compared with those with lower temperatures. Therefore, fever appears to have a favourable prognostic value in some diseases. The findings of our study suggest that most infants with bronchiolitis are afebrile (69%) and that fever is associated with more severe bronchiolitis.

In respiratory diseases, particularly those occurring in young infants, such as bronchiolitis, hypoxia is common and may be severe. In our series, 38 of the 90 children had severe bronchiolitis requiring supplemental oxygen to maintain adequate oxygen saturation. A rise in body temperature results in an increase in energy expenditure of about 10% for each 1°C rise in temperature. These changes are accompanied by an increase in oxygen consumption of 10–12% for every 1°C rise in temperature. This may explain why a high proportion of the febrile children (20 of 28) had a severe clinical course of bronchiolitis requiring oxygen supplementation. The necessity for oxygen supplementation was one of the main factors contributing to prolonged hospitalisation. It is possible that a reduction of body temperature by frequent use of antipyretic agents could reduce the oxygen requirement and possibly the degree of hypoxia.

Fever in respiratory diseases can increase the rate of pre-existing tachycardia and tachypnoea, which are caused by the disease itself. Therefore, tachycardia and tachypnoea may not be reliable signs of disease severity in febrile infants. Hence, oxygen saturation and the need for oxygen supplement were taken as criteria of disease severity. Furthermore, some infants with less severe bronchiolitis, who would not initially require oxygen supplementation, may do so later because of additional tachycardia and tachypnoea as a result of fever.

The presence of segmental/lobar collapse with consolidation in most of the febrile children might also have caused both a febrile response and a severe clinical course requiring an increase in oxygen consumption. Because many children with a severe clinical course were afebrile (29%) and had a normal chest x-ray, fever and radiological abnormalities do not seem to be the only factors contributing to clinical severity. Although some reports suggested a correlation between radiological abnormalities on a chest x-ray and disease severity, others could not confirm such a correlation. Fever, however, was not studied in these reports. Our study suggests that there is a correlation between disease severity and radiological changes in both febrile and afebrile children. Because febrile children had a higher incidence of such severity, more radiological abnormalities were expected in this group.

In contrast to other studies, which reported that children with bronchiolitis younger than 3 months of age have a more severe clinical course, in our study, the mean age of children with greater disease severity was higher than those with milder disease. Fever was, however, not included in these studies. In our study, children with a severe presentation within the same febrile or afebrile group were actually younger than those with a mild presentation (data not shown).

The reason why most of the 90 children were afebrile is not entirely clear. Production of interferon is enhanced as an important defence mechanism in many respiratory viral infections, including bronchiolitis. Interferon is known to be a potent endogenous pyrogen capable of inducing fever. However, the production of interferon by blood mononuclear cells has been shown to be significantly reduced during acute RSV bronchiolitis. This may explain the low incidence of fever in bronchiolitis. In addition, infants below the age of 3 months are less likely than older children to mount a febrile response, despite severe infection, because they have an immature thermoregulatory system and a lower metabolic rate than older children.

We thank Dr Kensing and his team from the microbiology department, Queen Mary’s Hospital, Sidcup, for their assistance in processing the nasopharyngeal aspirates from the studied children. We also thank the nursing staff on the Whiteoak Ward for their help during the study.

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