Clinical features, aetiology and outcome of empyema in children in the north east of England

K M Eastham, R Freeman, A M Kearns, G Eltringham, J Clark, J Leeming, D A Spencer

Background: The incidence of empyema in children in the UK is increasing. The reason for this is unclear. A prospective study was undertaken to investigate the clinical features, aetiology, and outcome of cases of empyema and parapneumonic effusion presenting to a tertiary paediatric respiratory centre between February 1997 and August 2001.

Method: Routine bacterial culture of blood and pleural fluid was performed for 47 cases. Forty three pleural fluid specimens, culture negative for pneumococcus, were analysed for pneumococcal DNA by real time polymerase chain reaction (PCR). Penicillin susceptibility was determined for DNA positive specimens using complementary PCR assay. Capsular serotype specific antigen detection was by enzyme immunoassay (EIA) using monoclonal antibodies to serotypes 1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F, and 23F. Clinical data were obtained from patient notes, supplemented by a postal questionnaire.

Results: The median (range) age of the patients was 5.6 (0.6–16.9) years and 70% were male. The median (range) duration of illness before referral to hospital was 5 (0–25) days. Forty five (96%) had received antibiotics before referral; 32 (68%) required decortication and eight (21%) thoracocentesis. Median postoperative stay was 4 days (2–8). Thirty two (75%) pneumococcal culture negative specimens were pneumococcal DNA positive; 17 (53%) of these were serotype 1. All were penicillin sensitive.

Conclusions: Pneumococcus is the major pathogen in childhood empyema and serotype 1 is the prevalent serotype. This has implications for vaccine development and immunisation strategy as the current 7-valent pneumococcal conjugate vaccine does not protect against serotype 1.

E mpyema and parapneumonic effusion complicating pneumonia has increased dramatically in children in the UK over the last decade. An explanation for this phenomenon has yet to be determined. Understanding of the problem has been hampered by the fact that routine bacterial culture is negative in most cases, presumably because antibiotics have usually been administered before referral to the regional centre. Empyema and complex parapneumonic effusion represent parts of a spectrum of disease, with three stages of progression being recognised:

- **Stage 1** ("exudative"): fluid accumulation within the pleural cavity without the presence of loculations (parapneumonic effusion).
- **Stage 2** ("fibropurulent"): fibrin deposition within the pleural space giving rise to loculations; presence of pus (empyema).
- **Stage 3** ("organisational"): organised multiloculated empyema with lung entrapment and pleural rind formation.

Management at our centre is determined by clinical features aided by ultrasound findings, as described previously.

*A streptococcus pneumoniae* has been found to be the principal pathogen in childhood empyema in the USA, with serotype 1 accounting for 24–50% of culture positive cases between 1993 and 2000. Clonal spread of this serotype has been proposed as a possible explanation. This is supported by the recent emergence of a novel clone of serotype 1 in Sweden which has been associated with a fourfold increase in invasive pneumococcal disease. In contrast, serotype 14 has been the most prevalent serotype causing invasive pneumococcal disease in England and Wales, although empyema related strains have not been specifically collected or typed.

The purpose of this investigation was to describe the clinical features, aetiology, and outcome of empyema and complex parapneumonic effusion in children presenting to Freeman Hospital, a tertiary paediatric centre in the north east of England, and to investigate the role of *S pneumoniae* using new microbiological techniques.

**METHOD**

Forty seven consecutive cases of empyema or complex parapneumonic effusion were recruited between February 1997 and August 2001. Referrals were all from other hospitals in the Northern Region and north Cumbria (Whitehaven and Carlisle). Informed parental consent was obtained. All cases were less than 17 years of age with clinical and radiological evidence of empyema or pleural effusion requiring management by thoracostentesis and/or decortication. Ultrasound examination staged the disease in 20 cases. The study was approved by the Newcastle and North Tyneside local research ethics committee.

A sample of 10 ml pleural fluid was collected at the time of chest drain insertion. Routine bacterial culture (including methods for anaerobes) and latex agglutination tests for pneumococcal antigen (SLIDEX, BioMérieux, France) were performed in all cases. In 43 cases sufficient pleural fluid was available for further analysis. A real time polymerase chain reaction (PCR) assay was performed for the detection of pneumococcal DNA. Pneumococcal DNA positive samples underwent a second complementary PCR assay targeting a conserved sequence of the *pbp2b* gene which is found in all strains of penicillin susceptible pneumococci (minimum inhibitory concentration to penicillin <0.06 mg/L). A solid phase indirect sandwich enzyme linked immunosorbent assay (ELISA) was performed on the 43 specimens to detect type specific pneumococcal capsular polysaccharide using
monoclonal antibodies against types 1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F and 23F (donated by Wyeth Vaccines).

Demographic and clinical data included age, sex, past medical history, duration of illness at presentation and referral, microbiological and radiological investigations, antibiotic use, length of hospital stay, and clinical outcome. Spirometric tests were performed at least 1 year after decortication in 11 of 16 children aged more than 5 years in May 2001. Data were collected by a postal questionnaire issued in October 2001 (3–56 months after admission to hospital) on work or school absence of family members during the acute illness.

RESULTS
Clinical features
Figure 1 illustrates case frequency, and the age distribution of the 47 children is shown in fig 2. The median age was 5.6 years (range 0.6–16.9); the male:female ratio was 2:1. Thirty one of the children (66%) had been previously fit and well; four had a concurrent infection at time of presentation (three varicella zoster and one Campylobacter gastroenteritis); eight had experienced a significant previous respiratory morbidity (three bronchiolitis, three recurrent respiratory tract infection, two asthma); two had a haematological malignancy and were undergoing chemotherapy; and two children had a chronic neurological disability with impaired mobility (spastic quadriplegia and metachromatic leucodystrophy).

The median duration of illness before admission to hospital was 5 days (range 0–25) and the median primary hospital stay was 3 days (range 0–14). Forty five cases (96%) had received antibiotics before transfer; 29 (61%) had received an antibiotic combination which included a cephalosporin and 22 (47%) included a penicillin.

Laboratory findings
A pathogen was implicated in 36 of the 47 cases (31 S pneumoniae; one dual infection with S pneumoniae and cytomegalovirus; one Staphylococcus aureus; one Streptococcus pyogenes; one Streptococcus viridans; and one Mycobacterium tuberculosis). All children had a blood culture taken at Freeman Hospital and six were positive (five S pneumoniae and one S viridans). All five cases with pneumococcal positive blood cultures were subsequently shown to have DNA evidence of pneumococcal infection in the pleural fluid. Routine bacterial culture of pleural fluid was negative in 44 cases (94%). Three specimens were culture positive: one Staphylococcus epidermidis which was considered to be a contaminant, one S pyogenes, and one S aureus in a child with a coexistent lung abscess. Mycobacterium tuberculosis (TB) was diagnosed by PCR in one child with an operative finding of pleural plaques in whom there was histological evidence of necrotising granulomata. Cytomegalovirus was isolated from viral culture of pleural fluid in a further child, performed because she had acute lymphoblastic leukaemia (ALL) and was undergoing chemotherapy.

Pneumococcal latex agglutination antigen testing was positive in 12 of 47 cases, all accounted for on PCR. Sufficient pleural fluid was available for further analysis in 43 of the 47 cases (fig 3). All were negative for S pneumoniae on routine culture. Pneumococcal DNA was identified by PCR in 32 cases (75%), illustrating the poor sensitivity of pneumococcal latex agglutination testing relative to PCR. Additional molecular analysis showed the presence of the conserved sequence associated with penicillin susceptibility in all 32 specimens. Figure 4 shows the serotype distribution.

Management
Surgical management formed the mainstay of treatment. Unsuspected lung abscess was an operative finding in seven cases. Pneumococcus serotype 1 was confirmed in three cases, serotype 14 in two cases, and serotype 3 in one case of lung abscess. Thirty two children (68%) required decortication including four cases managed initially by thoracocentesis. A further eight patients required thoracocentesis alone. Three children were managed initially with intrapleural urokinase but this failed to effect clinical resolution in two patients in whom decortication was subsequently required.

The median postoperative time to defervescence was less than 24 hours (range 0–6 days). Median stay in the intensive care unit for the cohort was less than 24 hours (range 1–48) and median postoperative inpatient stay was 4 days (range 2–8). Five children were found to be anaemic, three of whom were given a blood transfusion and two oral iron. Two children were found to have pre-renal failure secondary to hypovolaemia. Renal function was normal at discharge following appropriate fluid management. T cell lymphoblastic lymphoma was diagnosed in one child in whom a pleural effusion recurred 10 days after discharge.

Figure 1 Number of cases of childhood empyema and parapneumonic effusion presenting to Freeman Hospital, February 1997 to August 2001.

Figure 2 Age distribution of cases of empyema and parapneumonic effusion.

Pneumococcal DNA +ve (n = 32)
Pneumococcal DNA -ve (n = 13)
Penicillin sensitive (n = 32)

Figure 3 Detailed microbiological analysis of 43 pleural fluid specimens.
the USA has been attributed to pneumococcal serotype 1 in
had reproducible flow-volume loops. The median forced
(range 4–75) lost for 15 mothers, and 30 school or nursery
days (range 1–30) were lost for 17 fathers, 30 working days
beginning of onset of symptoms. A median of 10 working
enough to attend, was based on a 5 day week from the
and missed from school or nursery school for children old
for employed parents as a direct result of the child’s illness,
children, serotype 14 in one; and serotype 3 in one).
Serotyping was performed in five of these cases, confirming
pneumococcal titres was 2.4 years (range 1.6–6.4).
(range 70–108), median forced vital capacity (FVC) was 90%
voestalpine's performance with a bronchodilator.
Children with an obstructive defect, all of whom had evidence
persistence of the organism in infected tissue.
serotype 1 pneumococci have a characteristic that facilitates
increased prevalence of this serotype in the developing
and decreased availability of antibiotics have resulted in the
emerged in the community. Serotypes 1, 2 and 3 were
responsibility for 75% of bacteraemia in the western world in
emerged in the community. Serotypes 1, 2 and 3 were
pneumonias or that a virulent clone of pneumococcus has
argued that introduction of the vaccine has favoured the
serotype 1 in childhood empyema in the UK remain to be
August 2001 and July 2003. It is important to speculate why
this has occurred and why serotype 1 might be re-emerging
with such virulence in western societies.
The median age of our series was 5.6 years. This is
consistent with the US data16 and suggests that either type
1 pneumococcus has increased virulence in childhood
pneumonias or that a virulent clone of pneumococcus has
emerged in the community. Serotypes 1, 2 and 3 were
responsible for 75% of bacteraemia in the western world in
the early 20th century.15 16 Serotype 1 is seen now most
commonly in developing countries such as Rwanda and
Egypt.15 It has been suggested that poverty, overcrowding,
and decreased availability of antibiotics have resulted in the
increased prevalence of this serotype in the developing
world,16 although this does not explain its emergence in the
affluent state of Utah in the USA.7 Patients with pneumonia
complicated by a pleural effusion are more likely to be older,
of white race, and have chest pain on presentation than
patients with uncomplicated pneumonia.7 We propose that
serotype 1 pneumococci have a characteristic that facilitates
persistance of the organism in infected tissue.
Pneumococcal vaccination has been offered in the routine
immunisation schedule in the USA since the year 2000.17 The
7-valent pneumococcal vaccine protects against serotypes 4,
6B, 9V, 14, 18C, 19F, and 23F but not serotype 1. It could be
argued that introduction of the vaccine has favoured the
proliferation of novel clones not covered by it. However, this
could not account for the observations made in US studies6 7
in patients recruited before the introduction of widespread
vaccination or for our observations, as the 7-valent pneumo-
coccal conjugate vaccine has only recently been introduced in
high risk groups in the UK.14 Furthermore, the incidence of
complicated community acquired pneumonia has been
reported to have decreased since the introduction of the 7-
valent pneumococcal conjugate vaccine in the USA.12
Selection is unlikely to be driven by antibiotic resistance as
all the strains are penicillin sensitive (cephalosporins were
used most widely in our study population). We hypothesise
that the phenomenon of antibiotic tolerance, whereby
bacteria evade the killing action of the antibiotic without
an increase in the minimum inhibitory concentration (MIC),
may be a potential explanation for the re-emergence of
serotype 1 in western societies.20 Alternatively, the emerging
serotype may be producing an as yet unrecognised factor
important in its survival.
The factors responsible for the dominance of pneumococcal
serotype 1 in childhood empyema in the UK remain to be
determined. If antibiotic tolerance is found to be a

![Figure 4 Pneumococcal serotype distribution of culture negative cases.](http://www.thoraxjnl.com)
pathophysiological mechanism, this would have widespread implications for antibiotic policy in the community and hospital management of lower respiratory tract infections. Although a recent analysis of the cost of pneumococcal disease in childhood suggests that childhood empyema does not constitute a significant economic burden, we suggest that the severity and increasing prevalence of this condition does support a strategy of routine pneumococcal vaccination in early childhood. This would require the introduction of next generation vaccines (9-valent) to ensure the full impact in early childhood. This would require the introduction of next generation vaccines (9-valent) to ensure the full impact.

REFERENCES

Clinical features, aetiology and outcome of empyema in children in the north east of England

K M Eastham, R Freeman, A M Kearns, G Eltringham, J Clark, J Leeming and D A Spencer

Thorax 2004 59: 522-525
doi: 10.1136/thx.2003.016105

References

This article cites 21 articles, 11 of which you can access for free at:
http://thorax.bmj.com/content/59/6/522#BIBL

Email alerting service

Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Topic Collections

Articles on similar topics can be found in the following collections

- Inflammation (1020)
- Drugs: infectious diseases (968)
- Vaccination / immunisation (158)

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/