Empyema: the use of broad range 16S rDNA PCR for pathogen detection

S Saglani, K A Harris, C Wallis, J C Hartley

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Background: An increase in the incidence of thoracic empyema in children has been reported. The causative pathogen is often unknown as pleural fluid is frequently sterile at the time of culture. The role of unusual organisms is unclear.

Aims: (1) To compare the detection of organisms in pleural fluid from children with empyema using a molecular technique (16S rDNA polymerase chain reaction (PCR)) and bacterial culture. (2) To compare the concordance of organisms identified using the two techniques and the influence of prior antibiotic treatment on positive detection rate.

Methods: Pleural fluid from children admitted with empyema between January 2000 and February 2002 was cultured and additionally analysed using broad range 16S rDNA PCR.

Results: Pleural fluid was cultured from 32 patients, aged 1 month–16 years. Median duration of previous antibiotic therapy was 8 days (range 1–42 days). Six samples were culture positive and 22 were PCR positive. A causal organism was detected by PCR alone, after considering results from the local hospital, in 14 patients. There was complete concordance in organisms cultured and detected by PCR. Additional organisms detected by PCR were predominantly S pneumoniae, S pyogenes, and anaerobes.

Conclusions: Analysis of pleural fluid by broad range 16S rDNA PCR in addition to culture, increases organism identification in empyema.
instructions. One negative control (200 μl of sterile, UV irradiated water) was included in each extraction run.

**PCR amplification**

Five microlitres of extracted DNA and 5 μl of extracted DNA diluted 1/10 were amplified with primers 16SFa, 16SFb, and 16SR as previously described. 15

**Sequencing**

PCR products were sequenced with the ABI PRISM Big-dye sequencing kit (PE Applied Biosystems, Warrington, UK) and analysed on the ABI 377 Genetic Analyser (PE Applied Biosystems). Sequences were compared to the Genbank database using the BLAST program available at The National Centre for Biotechnology Information (www.ncbi.nlm.nih.gov) and to an in-house database of 16S rDNA sequences using the CLUSTAL algorithm in the MEGALIGN computer program (DNASTar, Madison, WI). Criteria for bacterial identification were as previously described. 15

**Assay sensitivity/specificity**

In silico, the 16S rDNA PCR can detect all common, known pathogens, including *Mycobacterium tuberculosis*. From control cultures of *Escherichia coli* and *Staphylococcus aureus* the assay can detect DNA 10–100 colony forming units/PCR reaction input. Precise details of assay performance and the genera of specific organisms that have been identified using the assay have been previously described. 15

**Statistical analysis**

In order to calculate the sensitivity and specificity of the broad range PCR in detecting organisms in empyema, we required a reference standard against which to compare the PCR results. 16 This standard is bacterial culture, but as so few samples were culture positive, this calculation was meaningless. Therefore the assay sensitivity could only be calculated based on the assumption that all patients should potentially have had organisms in their pleural fluid (and therefore should be culture positive), as they all had a clinical picture of empyema.

**RESULTS**

Thirty two patients were included, median age 2.9 years (range 1 month–16 years). Six patients (19%) had pleural fluid aspiration prior to transfer to our centre; four of these (67%) were culture positive. Patients had received a median of 8 days’ (range 1–42 days) antibiotic therapy prior to fluid aspiration at our centre (fig 1). They had all received antibiotics that would act against penicillin sensitive strains of *S pneumoniae*; however, details of previous anti-staphylococcal therapy were not known.

For samples collected in our centre, significant organisms were cultured in 6/32 (18.7%) cases (three *S aureus*, one *S milleri*, one *S pyogenes*, one *S pneumoniae*), whereas 22/32 (68.7%) cases were PCR positive (24 organisms identified). Pleural fluid was PCR positive and culture negative in 17/32 (53%) cases (table 1). Table 2 shows the organisms identified by PCR or culture in the 32 samples.

Four patients had a growth of coagulase negative staphylococci (three on subculture only), thought to be a contaminant growth. These were considered culture negative for the purposes of this study. Two of the four were PCR positive, both with *S pneumoniae*, one was PCR negative, and the remaining sample revealed a mixed sequence, felt to be contaminants, and was therefore reported negative.

Five of six culture positive samples were PCR positive; the organism isolated by culture was the same as that identified using PCR in all cases; the organism not detected by PCR was a *S aureus* grown only after enrichment culture. Six of 32 patients had previously had pleural fluid cultured at their local hospital. Four of these patients had a positive culture result from the local hospital. Pleural fluid samples from these cases were all culture negative, but three were PCR positive at our centre. The organisms identified by PCR were the same as those isolated at the local hospital (two *S pneumoniae*, one *S pyogenes*).

After considering all results (local and our centre) in the PCR group, causative organisms were detected in 24/32 (75%) cases (table 2). One case was only culture positive at the local hospital, and one only culture positive at our centre (table 2). An organism was identified by PCR alone in 14/32 (44%) patients.

The sensitivity of 16S rDNA PCR in this series of samples (for patients who had clinical empyema, but had received antibiotics was 22/32 (69%). The sensitivity of culture for the same samples was 6/32 (18.7%).

**DISCUSSION**

The overall rate of organism isolation using pleural fluid culture for our study was 18.7% (6/32), approximating previous reports, 8 17 and the range of organisms cultured were in close agreement with other studies from the UK. 18 Addition of broad range16S rDNA PCR to detect pathogens increased organism detection in pleural fluid collected at our centre from 18.7% (6/32) to 68.7% (22/32) of cases. In PCR positive samples from patients where organisms had been cultured in the referring hospital 1 or in a later specimen, there was complete concordance for bacterial identity, although such cases were small in number. Even taking into account results from samples collected and processed before referral to our unit, use of 16S rDNA PCR detected causal organisms in an additional 44% of children with empyema.

The main additional organisms identified by 16S rDNA PCR were *S pneumoniae* and *S pyogenes*. This is in agreement with the previously reported findings of a specific pneumococcal PCR being positive in the majority of a group of
children with empyema. However, DNA of more fastidious organisms was also detected by our broad range PCR. These were mainly anaerobic organisms, found in cases with an unusually protracted illness, which had shown a poor response to broad spectrum antibiotics. These organisms were detected using PCR, even though some of the children had the longest duration of antibiotic therapy (up to 42 days) prior to pleural fluid evaluation. The additional organisms detected by PCR were probably not cultured because of prior antibiotic treatment. S. pneumoniae and S. pyogenes are usually very sensitive to penicillins (common agents used to treat respiratory infection), and may have been rendered unculturable by prior treatment. S. aureus is usually resistant to penicillin and may therefore have persisted. In this study there was one case in which S. aureus was cultured, after enrichment, but not detected by PCR. This result occurred because the organism was present at a level below the detection limit of the PCR assay. Also, one sample was both culture and PCR negative at out centre, but had been positive at the referring hospital. This is because the period of time between antibiotic administration and empyema sampling alters the culture result, as organisms are killed, and the PCR result, as subsequent DNA degradation takes place. DNA remains after organisms become non-viable by culture methods, but DNA does not persist indefinitely and PCR subsequently also becomes negative. It is therefore important to emphasise that PCR should be complementary to culture, and not used as a replacement.

An increase in penicillin resistant S. pneumoniae may be contributory to the increase in the incidence of empyema. However, it is possible to determine whether the organisms identified by PCR alone were penicillin sensitive, using molecular assays based on the penicillin binding protein 2B gene. This approach has been applied to a series of empyema samples and to a number of samples from this series (unpublished data, KA Harris). Only fully sensitive S. pneumoniae were identified. This suggests that the increase in empyema rate is not predominantly due to penicillin resistant S. pneumoniae.

The failure of culture to detect causative organisms in the majority of empyema samples in this series is due to prior antibiotic treatment. This is important as empirical antibiotic selection is influenced by anticipated causative organisms and expected sensitivity. This information suggests there is no need for broad spectrum antibiotics but that appropriate dose and selection of agents able to penetrate the empyema, allied to drainage of pus, are more important in prevention or resolution of this infection. However, even with PCR, 25% of the series still did not have a bacterial diagnosis. This is most likely because of previous treatment with antibiotics and bacterial DNA degradation. However, there may be other, yet undescribed organisms that were not detected. Importantly, although it appears that the PCR only has a sensitivity of 69%, this is in samples from patients who have all received previous antibiotic therapy. As can be seen from fig 1, cases that were PCR negative had all received at least four days of prior antibiotic therapy. Therefore, molecular analysis of samples taken earlier in the illness (especially before prolonged antibiotic use) may further increase diagnostic yield. Also, the assay sensitivity could only be calculated based on the assumption that all pleural fluid samples should

### Table 2  Causative organisms detected in 32 cases of paediatric empyema, taking into account culture and PCR results from our centre and the local hospital

<table>
<thead>
<tr>
<th>Organisms</th>
<th>Method of detection</th>
<th>No. of patients</th>
<th>% of cases</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PCR</td>
<td>Culture</td>
<td></td>
</tr>
<tr>
<td>Streptococcus pneumoniae</td>
<td>13</td>
<td>1</td>
<td>14*</td>
</tr>
<tr>
<td>Streptococcus pyogenes</td>
<td>3</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>Staphylococcus aureus</td>
<td>2</td>
<td>3†</td>
<td>3</td>
</tr>
<tr>
<td>Fusobacterium sp</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Fusobacterium, Actinomyces, and</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Peptostreptococcus sp</td>
<td>1</td>
<td>1‡</td>
<td>1</td>
</tr>
<tr>
<td>Peptostreptococcus sp (PCR only)</td>
<td></td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>No bacterial cause detected</td>
<td>10</td>
<td>26</td>
<td>8</td>
</tr>
<tr>
<td>Total</td>
<td>32</td>
<td>32</td>
<td>32</td>
</tr>
</tbody>
</table>

*One positive culture from local hospital only.†One case was culture positive only at our centre.‡Only Streptococcus milleri group cultured.

### What this study adds

- First report of the use of a broad range PCR to detect causative organisms in a series of patients with empyema
- Use of the broad range PCR increases detection rate of causative organisms from 18.7% using culture alone, to 75% using both techniques
- There is complete concordance in organisms when identified both by culture and this molecular technique, but additional organisms are detected by PCR
- The overall pattern of causative organisms in empyema is unchanged and unlikely to be the explanation for the increasing incidence; however, some unusual organisms can be identified using this molecular technique

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have been culture positive, as culture is the reference standard against which to compare. However, culture only gave a sensitivity of 18.7%. Sensitivity has therefore been defined in samples of pleural empyema from children who have previously received more than one day of antibiotics. Earlier in the illness, or with less antibiotic treatment, the sensitivity of the PCR is likely to be higher.

In summary, this study has shown that the use of a molecular non-culture technique (broad range 16S rDNA PCR) improves organism detection rate from pleural fluid in children with empyema. Even after the commencement of antibiotics the causative organisms may be detected in 68.7% of cases. The majority of organisms detected were as expected, S pneumoniae, S pyogenes, S aureus, and anaerobes accounting for most cases. These results suggest that most of the increase in paediatric empyema is not accounted for by unusual organisms. We do not suggest that this technique should replace culture, but that it is a beneficial adjunct, especially after prior antibiotic therapy, and little clinical improvement. The assay can be offered as a routine service in the presence of unusual organisms. We do not suggest that this technique should replace culture, but that it is a beneficial adjunct, especially after prior antibiotic therapy, and little clinical improvement. The assay can be offered as a routine service in a specialist laboratory, and with a turnaround time of approximately 48 hours, a result can be obtained sufficiently quickly to allow appropriate alterations in management, whereby a change to more narrow spectrum antibiotics may be made soon into the patient’s in-patient stay.

Authors’ affiliations
S Saglani, C Wallis, Department of Respiratory Paediatrics, Great Ormond Street Hospital for Children, Great Ormond Street, London, UK
K A Harris, J C Hartley, Department of Microbiology, Great Ormond Street Hospital for Children, Great Ormond Street, London, UK

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