

Clinical and bacteriological profile of community-acquired pneumonia among children in and around Chengalpattu

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ABSTRACT

Introduction: Community-acquired pneumonia (CAP) has become the major cause of mortality among children despite the availability of potent antibiotics and effective vaccines. This created the interest to identify the most common bacteria causing pediatric CAP, which helps to reduce morbidity and mortality by early diagnosis and specific treatment. **Aims and Objectives:** To study the prevalence of CAP, most common organisms and its susceptibility pattern, risk factors, and clinical outcome of the disease. **Materials and Methods:** This was a hospital-based case series study which was carried out between April 2014 and July 2014. The study population included 52 children, in the age group of 3 months to 12 years, with clinical pneumonia. Blood samples, sputum, and gastric aspirates were collected and processed. Serology and antimicrobial susceptibility tests were also performed. Statistical analysis was performed with SPSS statistical software. **Results:** Children in the age group of 1-5 years were the most affected (46.15%) with male predominance (55.77%). The most common symptom was found to be fever (90.38%) followed by a cough (78.85%) and the most common organism isolated was *Streptococcus pneumoniae* (41.67%). The mortality was observed in two cases (3.84%). **Conclusion:** CAP is still the common cause of morbidity and mortality due to the virulence of pathogens. The resistance of methicillin-resistant *Staphylococcus aureus* strain to most of the antibiotics indicates the unnecessary use of antibiotics as one of the reasons for increasing resistance. Recurrent respiratory infections have been elicited as the risk factors for the development of severe illness.

Key words: Methicillin-resistant *Staphylococcus aureus*, Pediatric community-acquired pneumonia, *Streptococcus pneumoniae*

INTRODUCTION

Pediatric community-acquired pneumonia (CAP) is pneumonia acquired from the community in a previously healthy child and not been hospitalized within 14 days before the onset of symptoms.^[1]

CAP causes two million deaths in each year among the children aged <5 years.^[2] India has the greatest burden of this disease accounting for 43 million cases and 0.4 million deaths.^[3]

Bacteria are the most common cause of CAP, with *Streptococcus pneumoniae* isolated in nearly 50% of cases.^[4] Physicians need reliable data on the relative prevalence of different etiological events in patients'

area of residence, in addition to clinical, laboratory and radiological findings to initiate antibiotic treatment empirically.

The present study has been focused on identifying the most common bacteria causing pediatric CAP in this region that helps to reduce morbidity and mortality by early diagnosis and efficient treatment by specific antimicrobials.

Aims and objectives

To study the prevalence of CAP, most common organisms and its susceptibility pattern, risk factors, and clinical outcome of the disease.

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MATERIALS AND METHODS

The present case series study included 52 children in the age group of 3 months to 12 years with the symptoms of fever, cough, dyspnea, tachypnea, clinical signs, and radiological evidence of pneumonia. Babies in the perinatal period, children on anti-tuberculosis therapy, immunocompromised children and other pneumonias such as aspiration pneumonia, ventilator-associated and other hospital-acquired pneumonias, and lipoid pneumonia were excluded from the study.

Clinical samples such as expectorated sputum or induced sputum, gastric aspirate, and blood were collected from 52 children clinically diagnosed as pneumonia. The serum samples were also collected for the serological diagnosis of *Mycoplasma pneumoniae*. A detailed clinical examination was performed. In all cases, baseline investigations and chest X-ray were carried out.

The collected samples were processed and cultured on bacteriological media such as nutrient agar, MacConkey agar, blood agar, chocolate agar, and mannitol salt agar. The isolates were identified by Gram-stain, colony characteristics, and biochemical parameters.

After isolating and identifying the organisms, antibiotic susceptibility tests were performed as per CLSI guidelines using the Kirby Bauer's disc diffusion method. Cold agglutination test was performed with the patient's serum to detect infection with *M. pneumoniae*.

The disease pattern was assessed clinically, confirmed by radiological and microbiological methods. Clinical outcome was recorded, and the results were analyzed.

Statistical differences were analyzed using conventional Chi-square test, and $P < 0.05$ was considered significant. Data obtained was analyzed using SPSS statistical software.

RESULTS

The study group included 52 cases of clinical pneumonia. The disease pattern, outcome, and bacteria associated with pneumonia were investigated in this study.

Distribution of cases as per age and sex ($n = 52$)

The gastric aspirates collected from all the children were examined for acid-fast bacilli (AFB) by Ziehl-Neelsen technique and were found to be negative for AFB.

The serum samples were tested for *M. pneumoniae* by cold agglutination test showed a titer of $<1:16$. Hence, all the samples were found to be negative for antibody to *M. pneumoniae*.

DISCUSSION

In the present study, 52 children suspected of having clinical pneumonia were investigated. Pneumonia was radiologically confirmed in all the children. Most of the children ($n = 24$) with pneumonia were in the age group of 1-5 years (Figure 1) with a male:female ratio of 1.3:1 showing male predominance (Figure 2). Similar results were observed in a study carried out by Tullu *et al.*^[5] in which the male:female ratio was 2.4:1.

Fever was reported as the predominant symptom in 47 children (90.38%), followed by a cough (78.85%) and breathlessness (67.31%). Episodes of vomiting and diarrhea were also present in 9.61% and 5.77% of children, respectively (Figure 3). Campbell *et al.*^[6] also stated fever as a predominant and important clinical symptom of pneumonia.

In the present study, culture positivity rate of 23.08% was observed. The rate of isolation of the organism from sputum culture and blood culture was 13.46% and 9.61%, respectively (Table 1). Blood cultures are valuable when positive but negative results

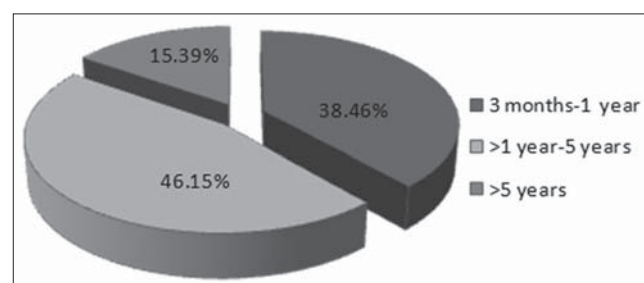


Figure 1: Age distribution. Chi-square value = 2.427; $P = 0.297$

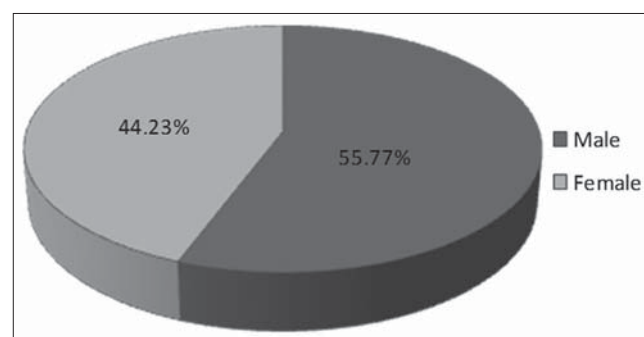


Figure 2: Sex distribution. Chi-square value = 0.028; $P = 0.867$

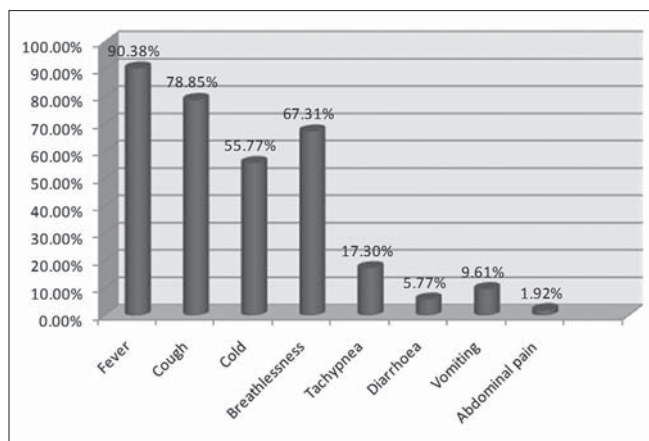


Figure 3: Clinical profile of children with pneumonia. Chi-square value = 1.372; $P = 0.927$ (for predominant symptom)

Table 1: Blood and sputum culture analysis (n=52)

Results	Number of cases	Percentage
Positive	12	23.08
Negative	40	76.92

are more common even in severe cases. Indian studies showed sputum culture positivity of 10-33%. In a study done by Falade *et al.*,^[7] blood culture was positive in 18% of cases. Agarwal^[8] reported blood culture positivity in 21.9% of cases. In our study, decreased culture positivity could be due to prior administration of antibiotics, and proper sputum collection is difficult in children.

The pathogen spectrum identified on culture was maximum number of *S. pneumonia* (41.67%), followed by MRSA (25%), Klebsiella pneumonia (16.67%), Staphylococcus aureus (8.33%), and coagulase-negative Staphylococci (CoNS) (8.33%) (Figure 4). Pneumococcal CAP is common in all age groups, and this is in well agreement with previous studies by Rao *et al.* from South India 2013^[9] and by Juvén *et al.* 2001^[10] and the other one from Texas USA by Wubbel *et al.* 1999.^[11] However, pneumococcal etiology was more frequent (42.10%) than the one found by Wubbel *et al.*^[11] (27%) and Juvén *et al.*^[10] (37%). In the present study, none of the cultures were found to be positive for Haemophilus influenzae Type b (Hib), probably because of fastidious growth requirements of the organism or due to the proper immunization schedules which include Hib vaccine as a part of pentavalent vaccine which is in accordance with the study by Karambelkar *et al.*^[12]

In the present study, antibiotic sensitivity pattern of *S. pneumonia* showed maximum sensitivity to penicillin, ampicillin, erythromycin,

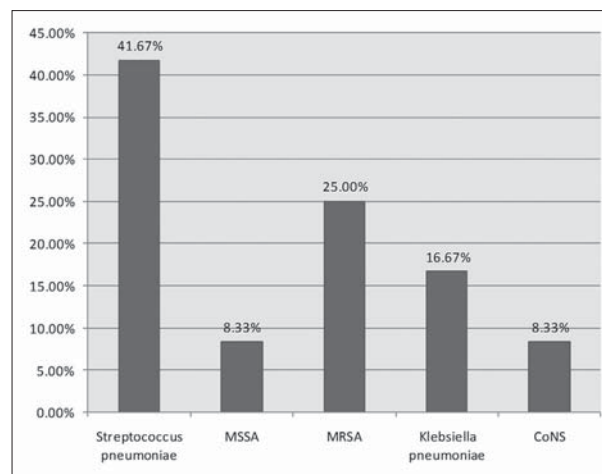


Figure 4: Organisms isolated by blood/sputum culture in children with pneumonia

cefotaxime, quinolones, and vancomycin. There were a total of four *S. aureus* isolated out of which three were methicillin resistant. The drug-resistant pattern of *S. aureus* to other drugs was as follows: Penicillin, ampicillin, erythromycin, gentamicin, cefotaxime - 75% each, amikacin - 50%, ciprofloxacin - 25%, and vancomycin - 0% (Figure 5). The IBIS study also reported that among *S. pneumoniae*, resistance to penicillin was rare (1.3%), and none of the isolates were resistant to injectable third generation cephalosporins.^[13,14]

In our study, none of the serum samples showed positive for *M. pneumonia*, the titer value was <1:16 in all the children. However, the studies done by Esposito *et al.*,^[15] *M. pneumonia* contributes to pediatric CAP (33.5%) and Principi and Esposito^[16] also stated that mycoplasmal infections were common in young children.

In our study, out of 52 children with clinical pneumonia, 50 (96.15%) recovered completely without any complications with standard line of treatment, including 13 children who had past history of similar illness, 3 known cases of developmental delay, 2 children with mental retardation and situs inverses totalis was found in 1 child.

In our study, mortality was observed (Table 2) in 2 cases (3.84%), with chest X-ray results as bronchopneumonia (1 case) and extensive pneumonitis (1 case). Out of these two cases, MRSA was isolated in one known case of developmental delay and methicillin-sensitive *S. aureus* (MSSA) in one, with chest wall deformity and respiratory distress. The study by Karambelkar *et al.*^[12] also showed mortality in CAP in children due to MSSA. The mortality due to

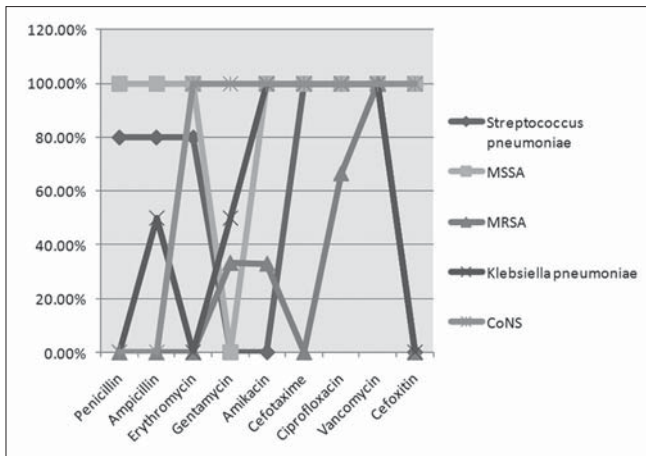


Figure 5: Sensitivity pattern of organisms isolated

Table 2: Analysis of clinical outcome/mortality in children with pneumonia (n=52)

Clinical outcome	Culture	
	Positive	Negative
Total number of cases	12	33
Death	2	0
Children well at time of discharge	10	40

Chi-square value=33.973; P=0.000

severe pneumonia among all hospitalized children was reported to be 1.35%, 3.32%, and 0.89% in a multi centric study carried out at Chandigarh, Kolkata, and Vellore.^[12] In the million death study done in India, among all the causes of childhood mortality, 27.6% deaths were attributed to pneumonia.^[17]

CONCLUSION

In the present study, more number of pneumonia cases in children were observed in the age group of 1-5 years with a male preponderance, and the most common symptom was fever followed by the cough and breathlessness. Blood culture and sputum cultures results were positive in 23.08% of cases, and the most common bacterial pathogen isolated were *S. pneumoniae* followed by MRSA, *K. pneumoniae*, MSSA, and CoNS.

The isolates showed a greater sensitivity to the antibiotics except the MRSA strain which was resistant to most of the antibiotics. The unnecessary use of antibiotics is one of the reasons for increasing resistance. Recurrent respiratory infections have been elicited as the risk factors for the development of severe illness. The mortality was observed in two cases, both the children were above 2 years of age and the bacterial pathogens isolated were MRSA and MSSA.

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REFERENCES

- Ghimire M, Bhattacharya SK, Narain JP. Pneumonia in South-East Asia Region: public health perspective. Indian J Med Res 2012;135:459-68.
- Rudan I, Tomaskovic L, Boschi-Pinto C, Campbell H; WHO Child Health Epidemiology Reference Group. Global estimate of the incidence of clinical pneumonia among children under five years of age. Bull World Health Organ 2004;82:895-903.
- Rudan I, Boschi-Pinto C, Biloglav Z, Mulholland K, Campbell H. Epidemiology and etiology of childhood pneumonia. Bull World Health Organ 2008;86:408-16.
- Shann F. Etiology of severe pneumonia in children in developing countries. Pediatr Infect Dis 1986;5:247-52.
- Tullu MS, Deshmukh CT, Baveja SM. Bacterial nosocomial pneumonia in paediatric intensive care unit. J Postgrad Med 2000;46:18-22.
- Campbell H, Byass P, Lamont AC, Forgie IM, O'Neill KP, Lloyd-Evans N, et al. Assessment of clinical criteria for identification of severe acute lower respiratory tract infections in children. Lancet 1989;1:297-9.
- Falade AG, Mulholland EK, Adegbola RA, Greenwood BM. Bacterial isolates from blood and lung aspirate cultures in Gambian children with lobar pneumonia. Ann Trop Paediatr 1997;17:315-9.
- Agarwal A. Bacteriological profile, serology and antibiotic sensitivity pattern of microorganisms from community acquired pneumonias. JK Sci 2006;8:79.
- Rao DR, Basu R, Sarkar A, Bidyarani K. Prevalence and antimicrobial susceptibility pattern of *Streptococcus pneumoniae* isolated from respiratory samples in a South Indian Tertiary Care Hospital. IJHSR 2013;3:121-6.
- Juvén T, Mertsola J, Toikka P, Virkki R, Leinonen M, Ruuskanen O. Clinical profile of serologically diagnosed pneumococcal pneumonia. Pediatr Infect Dis J 2001;20:1028-33.
- Wubbel L, Muniz L, Ahmed A, Trujillo M, Carubelli C, McCoig C, et al. Etiology and treatment of community-acquired pneumonia in ambulatory children. Pediatr Infect Dis J 1999;18:98-104.
- Karambelkar GR, Agarkhedkar SR, Karwa DS, Singhanian SS, Mane SV. Disease pattern and bacteriology of childhood pneumonia in Western India. Int J Pharm Biomed Sci 2012;3:177-80.
- The India Invasive Bacterial Infections Surveillance (IBIS) Study. International clinical epidemiology network. Prospective multicentre optical surveillance of Streptococcus pneumoniae in India. Lancet 1999;353:1216-21.
- Bansal A, Singhi SC, Jayashree S. Penicillin and gentamicin therapy vs. amoxicillin/clavulanate in severe hypoxemic pneumonia. Indian J Pediatr 2006;73:305-9.
- Esposito S, Bosis S, Cavagna R, Faelli N, Begliatti E, Marchisio P, et al. Characteristics of *Streptococcus pneumoniae* and atypical bacterial infections in

- children 2-5 years of age with community-acquired pneumonia. Clin Infect Dis 2002;35:1345-52.
16. Principi N, Esposito S. Emerging role of *Mycoplasma pneumoniae* and *Chlamydia pneumoniae* in paediatric respiratory-tract infections. Lancet Infect Dis 2001;1:334-44.
 17. Million Death Study Collaborators, Bassani DG, Kumar R, Awasthi S, Morris SK, Paul VK, *et al.* Causes of neonatal and child mortality in India: A nationally representative mortality survey. Lancet 2010;376:1853-60.
 18. Esposito S, Bosis S, Cavagna R, Faelli N, Begliatti E, Marchisio P, *et al.* Characteristics of *Streptococcus pneumoniae* and atypical bacterial infections in children 2-5 years of age with community-acquired pneumonia. Clin Infect Dis 2002;35:1345-52.

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