

Title	Community Acquired Pneumonia in Children
Document Type	Guideline
Issue no	CH024/02
Issue date	15/12/14, Reviewed September 2019
Review date	18/09/21
Distribution	Child Health
Prepared by	Graeme Eunson
Developed by	Dr Graeme Eunson
Equality & Diversity Impact Assessed	Yes

# Community acquired pneumonia (CAP) in childhood

CAP can be defined as the presence of signs and symptoms of pneumonia (fever, cough, tachypnoea) in a previously healthy child due to an infection acquired outside hospital.

## Key points:

1	Viral & bacterial pneumonia are indistinguishable clinically.
2	Children being discharged home do not need any investigations.
3	Antibiotics should be given if the diagnosis is clear, especially in children > 2yrs.
4	7 days of oral Amoxicillin is recommended, even in severe pneumonia.
5	IV antibiotics should only be used if oral will not be tolerated, or the child has septicaemia, empyema, abscess or necrosis.
6	If the child is feverish or unwell 48 hours after starting treatment, they must be re-evaluated for complications.

## Exceptions to the guideline:

- Infants < 6 months of age.
- Severe pneumonia.
- Pneumonia secondary to bronchiolitis.
- Underlying cardiac / respiratory condition or immunocompromise.
- Unwell child: eg shock, septicaemia.

### Clinical features:

- Fever, tachypnoea, difficulty breathing, cough, wheeze or chest pain.
- There are no distinguishing features between bacterial and viral pneumonia.
- Bacterial pneumonia should be considered in children with persistent fever > 38.5° with chest recession and tachypnoea.

#### Aetiology:

- Viruses account for about half of all cases of CAP, more in children < 2 years of age.
- One third of cases represent mixed infection (viruses + bacteria).
- Strep. pneumoniae = commonest bacterial cause.
- *Mycoplasma* is an important pathogen in all children (including pre-school).
- Group A streptococci & Staph. aureus are more likely than pneumococcal pneumonia to progress to empyema or ICU admission.

### Investigations:

- Children who do not require admission to hospital should have no investigations.
- CRP, WCC & CXR do not reliably distinguish between viral and bacterial pneumonia.
- Blood culture should be taken if IV therapy is required.
- Throat swab / NPA should not be done routinely in mild-moderate CAP.
- Microbiological diagnosis should be attempted only in severe CAP requiring ICU.

#### Severity assessment:

Mild-moderate	Severe
Temperature < 38.5	Temperature > 38.5
RR < 50	Tachycardia
Mild recession	RR > 70 in infants, > 50 in older children
Taking full feeds, no vomiting	Sats < 92%
	Severe difficulty breathing: recession, grunting, nasal flaring
	Apnoea, cyanosis
	Not feeding
	Signs of dehydration, CRT > 2 secs

### <u>Indications for admission to hospital:</u>

- Severe disease.
- Underlying risk factors eg prematurity, cardiac or respiratory conditions, immunocompromise.
- Ability of parents / carers to manage the illness at home, and distance of home from hospital.
- Weak cry and abnormal response to parental stimulation.
- Complications eg pleural effusion.

## **Management**

#### General measures:

- Antipyretics
- Oxygen to maintain sats > 92%.
- Fluids: enteral if tolerated, NG unless severely unwell. IV otherwise, with baseline & daily U&Es.
- Identify deterioration or other serious illness.
- How to access further healthcare & when to do so, ie safety net. Written or verbal information.

### Antibiotics: Decisions:

- 1. Whether to treat with antibiotics.
- 2. Which antibiotic and by which route.
- 3. When to change to oral if IV is started.
- 4. Duration of treatment.

Antibiotic resistance is uncommon in the UK & the clinical impact is not currently significant.

- Children with a clear diagnosis of pneumonia should receive antibiotics as it is not possible to distinguish between viral & bacterial causes.
- Children < 2 yrs with mild symptoms of LRTI should not receive antibiotics, but need to be reviewed if symptoms persist. Pneumococcal immunisation increases confidence in this decision.
- Amoxicillin is recommended as first choice for all children with pneumonia. It is effective against most bacterial pathogens, is well-tolerated & cheap.
- Alternatives are Co-amoxiclay, Cefaclor, Erythromycin, Azithromycin & Clarithromycin.
- Macrolides can be added at any age if there is no response to 1<sup>st</sup> line therapy.
- Macrolides should be used if Mycoplasma or Chlamydia pneumonia is suspected.
- In pneumonia associated with influenza, Co-amoxiclav is recommended.
- Oral antibiotics are safe and effective for all children, even in severe CAP.
- IV antibiotics should be only be used :
  - o when the child cannot tolerate oral fluids or absorption is in doubt (eg vomiting)
  - septicaemia
  - o complications of pneumonia eg empyema, abscess, necrosis

Amoxicillin is 1<sup>st</sup> choice. Co-amoxiclay, Cefotaxime, Cefuroxime, ceftriaxone are alternatives.

• There is no evidence for timing of IV to oral switch, consider when child is improving.

### Complications & failure to improve:

If child remains feverish or unwell after 48 hours, re-evaluation is necessary. Ask:

- Is the patient on appropriate drug treatment at an adequate dosage?
- Is there a lung complication eg empyema or abscess?
- Does the patient have co-existent disease eg cystic fibrosis or immunocompromise?

### Complications:

- Pleural effusions & empyema: ~1% of all CAP, up to 40% of those admitted to hospital.
- Necrotising pneumonia: predisposing factors: congenital cysts & sequestrations, bronchiectasis, neurological disorders & immunodeficiency, certain pneumococcal serotypes. CXR will raise suspicion of necrosis or abscess, & should be confirmed by CT. Prolonged IV antibiotics are required until fever settles.

- Septicaemia & metastatic infection: likely to require HDU/ICU care. Osteomyelitis or septic arthritis should be considered with staph. aureus infection.
- Haemolytic uraemic syndrome: consider if pallor, profound anaemia & anuria.
- Milder complications e.g. prolonged cough, abnormal chest shape & asthma have been reported.

## **Specific complications:**

- Staph aureus: pneumatoceles, occasionally leading to pneumothorax. Detection of S.aureus warrants immunological investigation.
- Mycoplasma: rashes are most common. Rarely Stevens-Johnson syndrome, haemolytic anaemia, polyarthritis, pancreatitis, hepatitis, pericarditis, myocarditis, encephalitis, aseptic meningitis & acute psychosis.
- Strep. pneumoniae: empyema, necrosis, abscesses.

## Indications for transfer to ICU:

- Respiratory failure requiring ventilatory support.
- Pneumonia complicated by septicaemia.
- Sats <92% in FiO2 > 60%.
- Rising RR& HR with severe respiratory distress & exhaustion.
- Recurrent apnoea or slow irregular breathing.

#### When should the child be reassessed?

Inform parents of features that suggest inadequate treatment & require medical review:

- Fever: high swinging or persistent fever (should start to settle within 48hrs of treatment).
- Effort of breathing: increased work of breathing and raised RR.
- Effect of breathing: the child is agitated & distressed.

In hospital: as above, plus vital signs.

- Agitation may be a sign of hypoxia.
- Look for signs of overwhelming infection & septicaemia.
- Examine for development of effusion / empyema, suggested by prolonged fever.

### Follow-up:

- Follow-up CXRs are not necessary in previously healthy children who are recovering well. They should be considered in those with round pneumonia, collapse or persistent symptoms.
- Children with severe pneumonia, empyema or lung abscess should be followed up until they have recovered completely and CXR is near normal due to risk of fibrosis & bronchiectasis

Dr Claire Hathorn Dr Graeme Eunson
SpR Paediatrics Consultant Paediatrician

July 2014

### Reference:

British Thoracic Society guideline for community acquired pneumonia in childhood. *Thorax* 2011;66(Supp 2):ii1-23.