Community-acquired pneumonia (non COVID-19)

Straight to the point of care
# Table of Contents

## Summary

## Basics
- Definition 4
- Epidemiology 4
- Aetiology 4
- Pathophysiology 5

## Prevention
- Primary prevention 6
- Secondary prevention 7

## Diagnosis
- Recommendations 8
- Case history 29
- Risk factors 30
- History & examination factors 31
- Diagnostic tests 36
- Differential diagnosis 45
- Diagnostic criteria 47

## Treatment
- Recommendations 50
- Treatment details overview 76
- Treatment options 78
- Emerging 109

## Follow up
- Recommendations 111
- Complications 113
- Prognosis 114

## Guidelines
- Diagnostic guidelines 115
- Treatment guidelines 116

## Online resources

## References

## Images

## Disclaimer
Patients with community-acquired pneumonia (CAP) typically present with symptoms and signs consistent with a lower respiratory tract infection (i.e., cough, dyspnoea, pleuritic chest pain, mucopurulent sputum, myalgia, fever) and no other explanation for the illness. Older people present more frequently with confusion or worsening of pre-existing conditions, and without chest signs or fever.

Diagnostic confirmation in all patients presenting to hospital requires evidence of consolidation (new shadowing that is not due to any other cause) on chest x-ray. A chest x-ray should not be requested routinely for patients managed in the community.

The CURB-65 mortality risk score (hospital setting) or CRB-65 severity score (community setting) are the basis, together with clinical judgement, for deciding whether to manage the patient in hospital or at home and determining appropriate therapy.

Initial treatment is with empirical antibiotics, following national/international guidelines and local epidemiology. In hospital, antibiotics should be administered within 4 hours of presentation.

Sputum and blood samples should be sent for culture in people with moderate- or high-severity CAP, ideally before antibiotics are started, and legionella and pneumococcal urine antigen testing should be considered.

Patients with oxygen saturation <94% (or <88% in patients at risk of CO₂ retention) should receive supplemental oxygen.

Sepsis should be considered whenever an acutely unwell person presents with likely infection, even if their temperature is normal. Local protocols (e.g., Sepsis Six or Surviving Sepsis Campaign 1 hour care bundle) should be followed for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.
Community-acquired pneumonia (non COVID-19)

Definition
Community-acquired pneumonia (CAP) is defined as pneumonia acquired outside hospital or healthcare facilities. Clinical diagnosis is based on a group of signs and symptoms related to lower respiratory tract infection with presence of fever >38°C (>100°F), cough, mucopurulent sputum, pleuritic chest pain, dyspnoea, and new focal chest signs on examination such as crackles or bronchial breathing. Older patients present more frequently with confusion or worsening of pre-existing conditions, and without chest signs or fever.[1] This topic focuses on the diagnosis and management of CAP in adults.

Epidemiology
Globally, lower respiratory tract infections are the most deadly infectious disease, resulting in 3 million deaths worldwide in 2016.[4] CAP is a serious health problem with high morbidity and mortality in all age groups worldwide, and is a major burden on healthcare resources.[5]

A literature review found that the overall annual incidence of CAP in Europe is between 1.07 and 1.2 per 1000 person-years and 1.54 and 1.7 per 1000 people.[6] The incidence of CAP increases with age to 14 per 1000 person-years in adults aged ≥65 years, and the incidence of CAP appears to be significantly higher in men than in women.[6] It is the fifth leading cause of mortality in Europe.[7] Estimates of mortality among patients range from 1% to 5% in outpatients, from 5.7% to 14% in general wards, and from 34% to 50% in the intensive care unit (especially in ventilated patients).[8] [9] Another study reported that mortality rates of CAP in Europe vary widely from country to country, ranging between <1% and 48%.[10] The mortality rate for pneumococcal pneumonia is about 5%, rising to between 6% and 30% in adults with associated bacteraemia.[11] [12]

In the US, the annual incidence of CAP has been estimated at 24.8 cases per 10,000 adults.[13] Pneumonia and influenza, when considered together, were the eighth leading cause of death (13.5 deaths per 100,000 population) and the leading infectious cause in the US in 2016.[14]

Aetiology
Streptococcus pneumoniae (the pneumococcus) is the most common causative pathogen of CAP across a range of severities and patient ages.[15] [16] [18] [19] However, other studies have found that influenza virus is the most common cause of CAP in adults.[7] [13] In Europe and the US, S pneumoniae accounts for about 30% to 35% of cases.[5] [16] [20] Other bacterial causes include Haemophilus influenzae, Staphylococcus aureus (including MRSA), group A streptococci, and Moxarella catarrhalis.

Atypical bacteria are also common causes, although they vary in frequency depending on the year and any epidemics.[19] [21] The incidence of atypical pathogens in community-acquired pneumonia is approximately 22% globally, but this varies with location.[22] The most commonly reported atypical bacteria are Mycoplasma pneumoniae, Chlamydia pneumoniae, and Legionella pneumophila. M pneumoniae accounts for up to 37% of CAP patients treated as outpatients and 10% of patients who are hospitalised.[5] [16] C pneumoniae accounts for 5% to 15% of cases of CAP,[23] and L pneumophila (especially serogroup 1) accounts for 2% to 6% of CAP in immunocompetent patients.[24] A systematic review found that Chlamydia psittaci was the causative organism in 1% of patients.[25] However, a Dutch study identified C psittaci by polymerase chain reaction (PCR) of sputum (when available) as a cause of CAP in 4.8% of cases.[26]
*Pseudomonas aeruginosa* may also be prevalent in patients with pneumonia, depending on the region; however, it is more common in hospital-acquired and ventilator-associated pneumonia compared with CAP. It accounted for 7.7% of all isolates in CAP in a systematic review in China.[27]

Respiratory viruses are reported in about 10% to 30% of immunocompetent adults hospitalised with CAP.[16] [28] [29] [30] Influenza virus A/B, respiratory syncytial virus, adenovirus, rhinovirus, and parainfluenza virus are the most common viral causes of CAP in immunocompetent adults. Newer pathogens reported to cause CAP include metapneumovirus and coronaviruses.[31] Detection of viral causes is increasing because of the use of PCR.

Polymicrobial aetiology in CAP varies from 5.7% to 13%, depending on the population and the microbiological diagnostic test used.[16] [29] [32]

**Pathophysiology**

Pneumonia develops subsequent to the invasion and overgrowth of a pathogenic micro-organism in the lung parenchyma, which overwhelms host defences and produces intra-alveolar exudates.[33]

The development and severity of pneumonia is a balance between pathogen factors (virulence, inoculum size) and host factors. The likely microbial causes of CAP differ according to a number of factors, including differences in local epidemiology, the setting (outpatients, hospitalised, or intensive care unit), severity of disease, and patient characteristics (e.g., sex, age, and comorbidities).[16]

Microbes that are present in the upper airways can enter the lower airways by microaspiration. Nevertheless, the defence mechanisms of the lungs (innate and acquired) keep the lower airways sterile. The development of pneumonia indicates a defect in host defences, exposure to a particularly virulent micro-organism, or a large inoculum size.

Impaired immune response (e.g., caused by HIV infection or advanced age) or dysfunction of defence mechanism (e.g., through current or passive smoking, COPD, or aspiration) leads to greater susceptibility to respiratory infections in patients.[6]

Pathogens can reach the lower respiratory tract by 4 mechanisms:

- Inhalation, a common route of entry for viral and atypical pneumonia in younger healthy patients. Infectious aerosols are inhaled into the respiratory tract of a susceptible person to initiate infection.
- Aspiration of oropharyngeal secretions into the trachea, the primary route through which pathogens enter the lower airways.
- Haematogenous spread from a localised infected site (e.g., right-sided endocarditis)[34]
- Direct extension from adjacent infected foci (e.g., tuberculosis can spread contiguously from the lymph nodes to the pericardium or the lung, albeit rarely).

There is a new theory that CAP may result from dysbiosis of the normal lung flora, rather than invasion of pathogenic micro-organisms in a sterile environment; however, this model requires further research.[35]
Primary prevention

Pneumonia prevention is focused on the pathogens that cause disease, through vaccination and by managing the risks associated with disease development.

The main means of prevention are pneumococcal and influenza vaccination of at-risk people and smoking cessation.[1]

1. **Pneumococcal and influenza vaccination** of at-risk individuals. Public Health England recommends:[61]
   - **Pneumococcal vaccination**
     - **Adults aged 65 or over and at-risk groups aged 2 years or over**: a single dose of 23-valent pneumococcal polysaccharide vaccine (PPV23). At-risk groups are those with:[62]
       - Asplenia or dysfunction of the spleen
       - Chronic respiratory, heart, kidney, or liver disease
       - Diabetes
       - Immunosuppression
       - Cochlear implants
       - Cerebrospinal fluid leaks
       - Occupational risk.
     - At-risk patients should be offered immunisation at every opportunity (for example, when immunising against influenza or at routine consultations), and especially at discharge from hospital.[62]

   - **Influenza vaccination**
     - **Adults aged 65 or over**, **people aged 6 months or over who are in at-risk groups, and children aged 2 to 17 who are not in at-risk groups who are part of the phased vaccination roll-out in the UK**: annual influenza vaccine with an age-appropriate formulation, provided they do not have a contraindication. At-risk groups are pregnant women and those with:[63]
       - Chronic respiratory, heart, kidney, liver, or neurological disease
       - Diabetes
       - Immunosuppression
       - Asplenia or dysfunction of the spleen
       - Class III obesity (BMI ≥40).

Further information on vaccines, vaccination procedures, special patient populations, and current vaccination schedules in the UK can be found in the ‘green book’[62] [63] [Public Health England: immunisation against infectious disease – the ‘green book’] and the latest Public Health England vaccination schedule.[61] [Public Health England: complete routine immunisation schedule]

Vaccination schedules vary by location; consult local guidance for recommendations.

2. **Smoking cessation**
   - Smoking cessation is important for all patients, but particularly for those at risk of pneumonia and influenza. Offer advice according to national smoking cessation guidelines.[1] [National Institute for Health and Care Excellence: stop smoking interventions and services]
• Cigarette smoking, both active and passive, is a recognised independent risk factor for CAP.[1]

Secondary prevention

For all patients with CAP who smoke, offer advice according to national smoking cessation guidelines. [National Institute for Health and Care Excellence: stop smoking interventions and services] Explain to patients how smoking impairs natural mechanisms for eliminating pathogens and debris.[1]
Recommendations

Urgent

Suspect CAP in patients with symptoms and signs of an acute lower respiratory tract infection and, in a hospital setting, with new radiographic shadowing (consolidation) for which there is no other explanation.[1] [64] [65]

Think# Could this be sepsis?# whenever an acutely unwell person presents with likely infection, even if their temperature is normal. Follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.[66] [67] [68]

- Remember that sepsis represents the severe, life-threatening end of infection.[69]
- Pneumonia is one of the main sources of infection in sepsis.[70]
- It is important to make an early, accurate diagnosis of infection and use antibiotics appropriately.

Urgent: in hospital

Prioritise a chest radiograph for all patients with suspected CAP within 4 hours of presentation to hospital.[1] [64]

Order blood tests including:[1]

- Oxygen saturations to guide supportive treatment
- Arterial blood gases in patients with SpO₂ <94%, those with a risk of hypercapnic ventilatory failure (CO₂ retention), and all patients with high-severity CAP (see our Management - recommendations section for more detail on assessing severity)
- Urea and electrolytes to inform severity
- Full blood count, liver function tests, and C-reactive protein to aid diagnosis and for baseline measurements.

Assess oxygen requirements. Prescribe oxygen if oxygen saturation <94% and maintain at target range.[1] In patients at risk of CO₂ retention prescribe oxygen if oxygen saturation <88%.[71]

- Target oxygen saturation range of:
  - 94% to 96% in acutely ill patients who are not at risk of hypercapnia[72]
  - 88% to 92% in patients at risk of hypercapnia.[71]

Assess mortality risk using the CURB-65 score and your clinical judgement for all patients with pneumonia confirmed by chest radiography (see our Management - full recommendations section for more information).[1] [64]

- Score of 3-5: high-severity.
  - Score of 3 or more: discuss with a senior colleague at the earliest opportunity and manage as high-severity pneumonia.[1] [64]
  - Score of 4 or 5: arrange emergency assessment by a critical care specialist.[1]
- Score of 2: manage as moderate-severity pneumonia.[1] [64]
- Score of 0 or 1: manage as low-severity pneumonia.[1] [64]

Send sputum and blood samples for culture in people with moderate- or high-severity CAP, ideally before starting antibiotics.[1] Consider legionella and pneumococcal urine antigen testing.[64]
**Measure observations** initially at least twice daily, and more frequently (e.g., every hour) in those admitted to a critical care unit (high-dependency unit or intensive care unit).[1]

**Urgent: in the community**

Perform a chest radiograph ONLY if you are unsure of the diagnosis and radiography will help you to manage the acute illness. [1] [64]

- For the majority of patients you should make a clinical diagnosis.[1] [64]
- **Consider pulse oximetry if working in an out-of-hours setting** to assess disease severity and oxygen requirement.[1]

**Assess mortality risk** using the CRB-65 score (see Management - full recommendations ) and your clinical judgement to inform severity. [1] [64]

- Score of 3 or more (high-severity): admit patient to hospital immediately . [1] [64]
- Score of 1 or 2 (moderate-severity): consider referral to hospital . [1]
- Score of 0 (low-severity): treat most patients at home. [1] [64]

**Key Recommendations**

**Presentation**

Patients with CAP typically present with symptoms and signs consistent with a lower respiratory tract infection (i.e., cough, dyspnoea, pleuritic chest pain, mucopurulent sputum, myalgia, fever) and no other explanation for the illness (e.g., sinusitis or asthma). [64]

Remember to **consider atypical presentations** (without obvious chest signs). For example:

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea[73]
- Legionella pneumonia may present as constitutional upset, diarrhoea, and confusion[73]
- Pneumocystis pneumonia in immunosuppressed people may present only as cough, dyspnoea, and marked hypoxia[73]
- **Older people frequently present with non-specific symptoms (especially confusion) and worsening of pre-existing conditions, and without chest signs or fever**[1]
- Some people present with acute confusional states.[73]

**Do not use clinical features alone** to predict the causative agent or to influence your initial choice of antibiotic.[1] [74]

**Risk stratification**

Determine whether the patient should be managed in hospital or at home using the CURB-65 mortality risk score (hospital setting) or CRB-65 score (community setting) (see our Management - recommendations section for more detail on risk stratification). [1] [64]

**Risk stratification in hospital** *
**Severity of CAP** | **CURB-65 score** | **Management decision**
--- | --- | ---
High | 4 or 5 | A range emergency assessment by a critical care specialist [1] [64]
3 | Discussed with senior colleague at the earliest opportunity and manage as high-severity pneumonia [1] [64]
Moderate | 2 | Consider for short-stay inpatient treatment or hospital-supervised outpatient treatment [1] [64]
Low | 0 or 1 | Consider for outpatient treatment [1] [64]

*All patients with CAP confirmed by chest radiograph.*

**Risk stratification in the community**

<table>
<thead>
<tr>
<th>Severity of CAP</th>
<th>CRB-65</th>
<th>Management decision</th>
</tr>
</thead>
<tbody>
<tr>
<td>High</td>
<td>3 or more</td>
<td>Admit to hospital immediately [1] [64]</td>
</tr>
<tr>
<td>Moderate</td>
<td>1 or 2</td>
<td>Consider hospital referral for assessment and treatment* [1] [64]</td>
</tr>
<tr>
<td>Low</td>
<td>0</td>
<td>Consider for treatment at home* [1] [64]</td>
</tr>
</tbody>
</table>

**Imaging**

Confirm the diagnosis by chest radiograph in all patients presenting to hospital with suspected CAP. [1] [64]

- A definitive diagnosis of CAP requires evidence of consolidation on chest x-ray. [75]

In community settings base the diagnosis on signs and symptoms of lower respiratory tract infection, focal chest signs, and illness severity. [1] [64]

**Further investigations**

Discuss with a senior colleague any patient who does not improve as expected. [1]

- Consider repeat chest radiograph, C-reactive protein, white cell count, and further specimens for microbiology in patients not progressing satisfactorily after 3 days of treatment. [1]
- Consider referral to a respiratory physician. [1]

In patients with high-severity CAP who do not respond to beta-lactam antibiotics or in whom you suspect an atypical or viral pathogen, order polymerase chain reaction (or other antigen detection test) of sputum or other respiratory tract sample. [1]
Community-acquired pneumonia (non COVID-19)

Diagnosis

• Consider initial and follow-up viral and atypical pathogen serology.[1]

In the community consider assessing oxygenation via pulse oximetry, if available.[1]

• Oxygen saturation $<94\%$ is an adverse prognostic factor in CAP and also usually an indication for oxygen therapy.[76]

• Consider urgent hospital referral in these patients.[1]

General investigations are not necessary in the majority of patients presenting in the community,[1] but you should consider a point-of-care C-reactive protein test if you cannot make a diagnosis of CAP from the clinical assessment and you are not sure whether antibiotics are indicated.[64]

Full Recommendations

Clinical presentation

No constellation of signs and symptoms is predictive of CAP. However, patients typically present with:[1] [64] [65]

• Symptoms and signs consistent with a lower respiratory tract infection:
  • Cough and at least one other respiratory tract symptom:
    • Shortness of breath – usually present[1]
    • Pleuritic chest pain
    • Tachypnoea
    • Mucopurulent sputum – associated with bacterial pneumonia (scant or watery sputum is associated with an atypical pathogen)[1]
  • At least one systemic feature:
    • Rigors or night sweats – usually present, but less common in older patients[1]
    • Fever ($>38\,^\circ\text{C}$ [$>100\,^\circ\text{F}$]) – older patients may be afebrile[1]
    • Non-specific symptoms – may include myalgia, lethargy, malaise, anorexia. Confusion is often seen in older patients[1]
  • New focal chest signs on examination such as crackles or bronchial breathing
  • No other explanation for the illness.

Remember to consider atypical presentations of CAP (i.e., without obvious chest signs). For example:

• Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.[73]
• Legionella pneumonia may present as constitutional upset, diarrhoea, and confusion.[73]
• Pneumocystis pneumonia in the immunosuppressed may present only with cough, dyspnoea, and marked hypoxia.[73]
• Older people frequently present with non-specific symptoms (e.g., lethargy, malaise, anorexia, confusion) and worsening of pre-existing conditions, and without chest signs or fever.[1]
• Some people present with acute confusional states.[73]

Consider speed of symptom onset in your differential diagnosis:

• Symptoms developing within minutes may be suggestive of pulmonary embolism, pneumothorax, or a cardiac aetiology.

Practical tip

Think" Could this be sepsis?" whenever an acutely unwell person presents with likely infection, even if their temperature is normal. Follow your local protocol (e.g., Sepsis Six or
Surviving Sepsis Campaign 1-hour care bundle) for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.  

- Remember that sepsis represents the severe, life-threatening end of infection.
- Pneumonia is one of the main sources of infection in sepsis.
- It is important to make an early, accurate diagnosis of infection and use antibiotics appropriately.

Some symptoms and signs are more common with specific pathogens.

- Do not, however, use clinical features alone to predict the causative agent or to influence your initial choice of antibiotic.

Typical pathogens causing CAP in adults in the UK and their most commonly associated features
<table>
<thead>
<tr>
<th>Pathogens</th>
<th>Most commonly associated clinical features</th>
<th>Other features</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Streptococcus pneumoniae</em></td>
<td>Acute onset, high fever, and pleuritic chest pain</td>
<td><strong>Most common overall pathogen</strong>&lt;br&gt;More likely in the presence of cardiovascular comorbidity, increasing age&lt;br&gt;Bacteraemic <em>S pneumoniae</em> is more likely in:&lt;br&gt;• Females&lt;br&gt;• People with a history of excess alcohol intake&lt;br&gt;• People with diabetes mellitus&lt;br&gt;• People with COPD</td>
</tr>
<tr>
<td><em>Haemophilus influenzae</em></td>
<td>No specific defining features</td>
<td>More likely in older people and those with COPD</td>
</tr>
<tr>
<td><em>Legionella pneumophila</em></td>
<td>Diarrhoea, encephalopathy and other neurological symptoms, severe infection more likely and evidence of multisystem involvement (e.g., abnormal liver function tests, elevated serum creatine kinase)</td>
<td>More likely in young patients without comorbidities, smokers, immunocompromised people, people exposed to contaminated artificial water systems (e.g., air conditioning units, spas, fountains, repair of domestic plumbing systems)&lt;br&gt;<strong>Higher frequency in severe illness (patients in the intensive care unit)</strong>&lt;br&gt;Enquire about foreign travel</td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em></td>
<td>No specific defining features</td>
<td>More likely if preceding or concurrent influenza infection&lt;br&gt;<strong>Higher frequency in severe illness (patients in the intensive care unit)</strong>&lt;br&gt;Enquire about influenza symptoms as they are of predictive value. Influenza virus infection can be</td>
</tr>
</tbody>
</table>
### Diagnosis

<table>
<thead>
<tr>
<th>Pathogens</th>
<th>Most commonly associated clinical features</th>
<th>Other features</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>complicated by co-/secondary infection with \textit{S} \textit{aureus}</td>
</tr>
</tbody>
</table>

#### Atypical pathogens causing CAP in adults in the UK and their most commonly associated features \cite{1}

<table>
<thead>
<tr>
<th>Pathogens</th>
<th>Most commonly associated clinical features</th>
<th>Other features</th>
</tr>
</thead>
<tbody>
<tr>
<td>\textit{Mycoplasma pneumoniae}</td>
<td>In young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea\cite{73}</td>
<td>More likely in younger patients</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Epidemic years occur in roughly 4-yearly cycles</td>
</tr>
<tr>
<td>\textit{Chlamydia pneumoniae}</td>
<td>No specific defining features</td>
<td>None</td>
</tr>
<tr>
<td>\textit{Coxiella burnetii}</td>
<td>History of dry cough and high fever</td>
<td>More likely in males</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Exposure to infected animal sources (especially during parturition) is the main epidemiological link\cite{77}</td>
</tr>
<tr>
<td>\textit{Klebsiella pneumoniae}</td>
<td>No specific defining features</td>
<td>People with alcohol dependency are at higher risk of bacteraemic and fatal \textit{Klebsiella} pneumonia</td>
</tr>
</tbody>
</table>

#### History

Your history should cover risk factors to help you assess whether the patient has CAP, a lower respiratory tract infection, or an alternative diagnosis.\# You should also identify factors that may influence the management plan if CAP is diagnosed.

Be aware that you cannot make a definitive diagnosis of CAP from the history alone.

#### Medical history

(*Denotes a strong risk factor for CAP.*)

- Respiratory chronic diseases:
  - \textbf{COPD*, asthma, and bronchitis} are associated with a 2-fold to 4-fold increased risk of CAP\cite{6}
  - COPD is an independent risk factor for mortality in patients with CAP.\cite{39}
- Other chronic comorbidities:
  - \textbf{Chronic heart disease} \cite{6} \cite{39}
  - \textbf{Diabetes} \cite{6} \cite{39} – the risk of severe pneumococcal bacteraemia is higher in people with diabetes.\cite{55}
Community-acquired pneumonia (non COVID-19)

**Diagnosis**

**Practical tip**

Consider aspiration pneumonitis/pneumonia in patients at high risk of aspiration, such as those with chronic swallowing difficulties, those with organic neurological conditions (e.g., Parkinson’s disease, stroke, dementia), or those who cannot protect their airways easily.[1]

**Social history**

- Age ≥65 years*
  - Incidence of CAP increases significantly with age. Advanced age is associated with a higher mortality from CAP.[10]
- Residence in a nursing home*
  - Mortality rates due to pneumonia in nursing home residents have been reported to reach 55%. [78] [79]
  - Nursing home residents also have an increased risk of aspiration pneumonia.[80]
- Contact with children*
  - Regular contact with children is associated with an increased risk of CAP.[50]

**Lifestyle history**

- Alcohol use/misuse*
  - People who consume alcohol at all or in higher amounts have an 83% higher risk of CAP compared with people who consume no alcohol or lower amounts (relative risk of 1.83).[44] For every 10 to 20 g higher alcohol intake per day, there is an 8% increase in the risk of CAP.[44]
- Smoking*
  - Smoking is an independent risk factor for developing CAP.[81]
  - Passive smoking at home is also a risk factor for CAP in people aged 65 years or older.[43] [81]
- Poor oral hygiene
  - Poor oral hygiene (particularly dental dysaesthesia and wearing dental prostheses) may contribute to a higher risk of CAP in adults.[82]

**Drug history**

- Proton pump inhibitors, H2 antagonists, and prescribed opioids (particularly immunosuppressive opioids) are associated with CAP.[60]

**Physical examination**

Carry out a thorough examination, particularly of the **cardio-respiratory system**, to identify features consistent with CAP.

Check for:

- Fever (>38ºC [>100ºF])
- Raised respiratory rate
- Tachycardia
• **Focal chest signs** – none, some, or all of these may be present:
  - Crackles
  - Bronchial breathing
  - Decreased chest expansion
  - Dullness to percussion (suggests consolidation and/or pleural effusion)
  - Decreased entry of air.

Focus in addition on other areas (e.g., throat, legs) if the presentation suggests an alternative diagnosis, such as an upper respiratory tract infection, deep vein thrombosis, or pulmonary embolism.

**Imaging**

**In hospital**

**All patients on presentation**

Send all patients seen in hospital with suspected CAP for a chest radiograph as soon as possible to confirm the diagnosis, and within 4 hours of presentation to hospital. [1] [64]

- New shadowing (consolidation) on chest x-ray confirms the diagnosis of CAP. [1] [64]
- Reassess the patient if the chest x-ray shows no consolidation. [1]

---

*A high-quality chest radiograph is important to ensure accurate diagnosis and to avoid inappropriate antibiotic prescribing. One study reported that 29% of hospitalised patients treated for CAP did not have radiographic abnormalities.* [83]
Bear in mind that it is more difficult to obtain a high-resolution image from a person with class III obesity (body mass index ≥40).

Reserved for specific circumstances

Consider a chest computed tomography (CT) scan if the radiograph is of poor quality or an ill-defined consolidation is present. [1]

Consider a chest CT scan or other imaging investigations for 'complicated' pneumonia or atypical changes on a chest radiograph, such as cavitation, multifocal consolidation pattern, or pleural effusion. [84]

Chest radiograph showing left upper lobe cavitating pneumonia
From the collection of Dr Jonathan Bennett. Used with permission
Community-acquired pneumonia (non COVID-19)

**Diagnosis**

Left-sided pleural effusion

*From the collection of Dr R Light. Used with permission*
Increased opacification of the right perihilar region and superior segment of the right lower and upper lobes consistent with worsening aspiration pneumonia

From the collection of Dr Roy Hammond. Used with permission
### Community-acquired pneumonia (non COVID-19)

#### Diagnosis

<table>
<thead>
<tr>
<th>X-ray findings</th>
<th>Further imaging</th>
<th>Consider alternative diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Cavitation</strong></td>
<td>Chest CT scan</td>
<td>• Tuberculosis</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Lung cancer</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Pulmonary infarct</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Septic pulmonary emboli</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Infected bulla</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Lung abscess</td>
</tr>
<tr>
<td><strong>Multifocal consolidation</strong></td>
<td>Chest CT scan</td>
<td>• Staphylococcal infection</td>
</tr>
<tr>
<td>(note that it is the multifocal</td>
<td></td>
<td>• Tuberculosis</td>
</tr>
<tr>
<td>nature, not the consolidation</td>
<td></td>
<td>• Aspiration pneumonia</td>
</tr>
<tr>
<td>that is the distinguishing</td>
<td></td>
<td>• Allergic bronchopulmonary aspergillosis</td>
</tr>
<tr>
<td>feature)</td>
<td></td>
<td>• Cryptogenic organising pneumonia</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Drug hypersensitivity reaction</td>
</tr>
<tr>
<td><strong>Pleural effusion</strong></td>
<td>Chest ultrasound +/- guided</td>
<td>• Parapneumonic effusion</td>
</tr>
<tr>
<td></td>
<td>aspiration +/- chest CT scan</td>
<td>• Congestive heart failure</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Tuberculosis</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Pulmonary embolism</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Lung cancer</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Fungal infection</td>
</tr>
</tbody>
</table>

#### Practical tip

‘Complicated’ pneumonia refers to pneumonia that is complicated by the presence of parapneumonic effusion (an exudative pleural effusion associated with pulmonary infection), empyema (pus in the pleural space), abscess, pneumothorax, necrotising pneumonia, or bronchopleural fistula.

Around 40% of people who are hospitalised for pneumonia develop parapneumonic effusion.[85] Empyema is a type of pleural effusion that is difficult to distinguish from a parapneumonic effusion on chest radiograph.

Findings on a CT scan suggestive of a parapneumonic effusion (as opposed to empyema) include:[86]

- Usually small volume
- Normal meniscus sign
- Dependent
- No loculation.

‘Split pleura sign’ is not typical and is more specific for empyema.

Consider **CT pulmonary angiography** (CTPA) to rule out pulmonary embolism if symptoms came on quickly (within minutes) or pain and breathlessness preceded infective symptoms.[87]
In the community

Do not request a chest radiograph for patients with suspected CAP seen in the community unless:

- There is diagnostic doubt
- The patient is deemed to be at risk of underlying lung pathology (e.g., they have risk factors for lung cancer)
- Progress following treatment is not satisfactory at review.

In community settings base the diagnosis on signs and symptoms of lower respiratory tract infection, focal chest signs, and illness severity.[1] [64]

Debate: Ultrasound in the diagnosis of CAP

Although a chest radiograph showing new shadowing that cannot be attributed to any other cause is the ‘gold-standard’ for the diagnosis of pneumonia, it may not always be feasible in a community setting and it involves exposure to radiation.

- Emerging evidence has shown that lung ultrasound is a possible accurate diagnostic test for people with CAP. However, the benefits of its use in practice over chest radiography are still unclear.
- A meta-analysis of 12 studies looking at the diagnostic accuracy of lung ultrasound in people with CAP found a sensitivity and specificity of 0.88 and 0.86, respectively.[88] However, there were limitations, such as the large variability in the findings and the lack of heterogeneity of the studies reviewed.
- Further evidence is required before recommendations can be made.

Other investigations

General investigations

In a hospital setting

Arrange the following tests for all patients admitted to hospital.

Start with oxygen saturations and urea (and electrolytes) as these will inform supportive treatment and severity, respectively:[1]

- Pulse oximetry (preferably while breathing air) to assess oxygenation saturations
  - Oxygen saturation <94% is an adverse prognostic factor in patients with CAP and may be an indication for oxygen therapy[76]
- Arterial blood gas (ABG) measurements in patients receiving oxygen therapy [89]
  - Measure ABG in patients with SpO₂ <94%, those with a risk of hypercapnic ventilatory failure (CO₂ retention), and all patients with high-severity CAP [1]
- Urea (and electrolytes) to inform severity of disease
  - Urea >7 mmol/L (>19.6 mg/dL) counts for 1 point in the 6-point CURB-65 score to assess severity[1]
  - Chronic renal failure is a significant risk factor for mortality in patients with CAP.[57]
Always record the inspired oxygen concentration clearly as this is essential for interpreting blood gas results.

Order **full blood count, C-reactive protein, and liver function tests** to help identify underlying or associated pathology, and for baseline measurements.[1]

- **Full blood count**. Leukocytosis is often seen in people with CAP:
  - A white cell count of $>15 \times 10^9$ /L indicates a bacterial (particularly pneumococcal) aetiology.[1]
- **C-reactive protein (CRP)** to help rule out other acute respiratory illnesses and as a baseline measure:
  - A level $>100$ mg/L makes pneumonia likely[90]
  - A level $<20$ mg/L with symptoms for more than 24 hours makes the presence of pneumonia highly unlikely[90]
  - A failure of CRP to fall by 50% or more at day 4 is associated with an increased risk for 30-day mortality, need for mechanical ventilation and/or inotropic support, and complications.[91]
- **Liver function tests** to assess liver function:
  - Abnormal in patients with underlying liver disease and legionella infection[1]
  - Chronic liver disease is a risk factor for pulmonary complications in patients hospitalised due to pneumococcal pneumonia.[59]

Perform early **thoracocentesis** in all patients with pleural effusion as this can reveal an infected pleural space consistent with a parapneumonic effusion or empyema.[1]

- Drain pleural fluid in patients with an empyema or clear pleural fluid with pH $<7.2$.[1]

**Monitoring**

Measure observations initially at least twice daily, and more frequently (e.g., every hour) in patients admitted to a critical care unit (high-dependency unit or intensive care unit).[1]

- Pulse
- Blood pressure
- Respiratory rate
- Temperature
- Blood pressure
- Oxygen saturation (with a recording of the inspired oxygen saturation at the same time)
- Mental status.

All patients with high-severity CAP (high risk of death) should be reviewed at least every 12 hours until improvement. [1] #This should be done by a senior colleague and the medical team. [1]

**In the community**

General investigations are not necessary for the majority of patients with CAP who are managed in the community.[1] However, you should **consider a point-of-care CRP** if you cannot make a diagnosis of CAP from the clinical assessment and it is not clear whether antibiotics should be prescribed.[64]
Consider **assessing oxygenation via pulse oximetry** if available (e.g., if you are working in an out-of-hours setting).[1]

- Oxygen saturation <94% is an adverse prognostic factor in CAP and also may be an indication for oxygen therapy.[76]
- Consider urgent hospital referral in these patients. [1]

### Microbiological investigations

Be aware that the extent of microbiological testing in an individual patient is guided by severity, presence of risk factors (e.g., COPD), and disease outbreaks (e.g., legionella pneumonia).[1]

### In hospital

**Do not perform microbiological tests routinely in patients with low-severity CAP presenting in hospital.**[1] Empirical antibiotic therapy is associated with a good prognosis in these patients.[1]

### Blood cultures

**Order blood cultures, ideally before antibiotics are given, in all patients with moderate- or high-severity CAP** (as determined by the CURB-65 score – see our Management recommendations section).[1] [64]

- Isolation of bacteria can be highly specific in determining the microbial aetiology in people with moderate or severe CAP.[1] [64]
- Bacteraemia is also a marker of illness severity. However, many patients with CAP do not have associated bacteraemia.[1] Microbial causes of CAP that can be associated with bacteraemia include:[1]
  - *Streptococcus pneumoniae* (identified in around 60% of positive blood cultures)[92]
  - *Haemophilus influenzae* (identified in 2% to 13% of positive cultures)[92]
  - *Staphylococcus aureus* and *Klebsiella pneumoniae*.

**Do not order blood cultures in patients with confirmed CAP who have low-severity disease and no comorbid conditions.**[1]

#### Debate: Blood cultures

*There is debate around the practicality of ordering routine blood cultures in patients hospitalised with CAP. This is mainly due to low sensitivity, cost, and the fact that results rarely influence antimicrobial management.*

- In a study of 355 patients admitted to hospital for CAP, the proportion of false-positive blood cultures was 10%, and the proportion of true positives was 9% (95% CI, 3.3% to 5.5%).[93]
- Antibiotic therapy was changed on the basis of blood culture results in only 5% of patients (95% CI, 3% to 8%).[93]
- However, despite these limitations, most experts still recommend blood cultures in patients with high-severity CAP.[1]

### Sputum cultures

**Send sputum cultures in:**

- **All patients with moderate- or high-severity CAP** (as determined by CURB-65 score).[64]
  The British Thoracic Society recommends sputum cultures in patients with moderate-severity CAP only if they have not received prior antibiotic therapy [1]
Gram stain of sputum cultures

Order Gram stain of sputum cultures in patients with high-severity CAP or complications if available in your local laboratory.[1]

- Gram stain is an immediate indicator of the likely pathogen and can help with interpreting culture results.[1]

**Evidence: sputum Gram stain**

_A prospective study of 1390 patients with bacteraemic CAP found a sensitivity for sputum Gram stain of:[94]

- 82% for pneumococcal pneumonia
- 79% for _H influenzae_ pneumonia
- 76% for staphylococcal pneumonia.

Specificities ranged from 93% to 96%.

**Practical tip**

Carrying out routine sputum Gram stains for all patients is unnecessary.[1] The test has a low sensitivity and specificity, and often does not contribute to initial management. Problems include:[1]

- Patients may not be able to produce good specimens
- Prior exposure to antibiotics
- Delays in transport and processing of samples, which reduces the yield of bacterial pathogens
- Difficulty interpreting the results due to contamination of the sample by upper respiratory tract flora, which may include potential pathogens such as _S pneumoniae_ and ‘coliforms’ (especially in patients already given antibiotics).

Urine antigen testing

**Streptococcus pneumoniae**

Consider pneumococcal urine antigen tests for people with moderate- or high-severity CAP. [64]

- Urine antigen testing is useful for diagnosing pneumococcal pneumonia in adults and is less affected than blood/sputum cultures by prior antibiotic therapy.[1]

**Evidence: Urine antigen testing for pneumococcal pneumonia**

_Studies have shown significantly greater sensitivity rates for the pneumococcal urine antigen test than for routine blood or sputum cultures._ [95]

- Results remain positive in 80% to 90% of patients for up to 7 days after starting antimicrobial treatment.[95]

**Debate: Patient groups for pneumococcal testing**

_The British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE) guidelines differ in their recommendations regarding who should be tested for pneumococcal pneumonia using urine antigen._#
Community-acquired pneumonia (non COVID-19)

**Diagnosis**

- The BTS recommends testing all patients with high-severity CAP,[1] whereas NICE recommends considering testing in patients with moderate- or high-severity CAP.[64]
- A later comparison by the BTS of the key recommendations in the two guidelines (BTS published in 2009 and NICE in 2014) concluded that there were no major differences between them and, where there were differences, clinicians should follow the NICE guideline instead of the BTS guideline.[96]

**Legionella**

Order *legionella* urine antigen tests in all patients with specific risk factors and for all patients with CAP during outbreaks. [1] Consider testing also for people with moderate- or high-severity CAP.[64]

- It is important that *legionella* pneumonia is diagnosed promptly as it is associated with significant mortality and has public health significance.[1]
- Detection of *Legionella pneumophila* urinary antigen by enzyme immunoassay allows for rapid results early in the illness.[1]
- Legionella antigen testing by enzyme immunoassay is highly specific (>95%) and sensitive (80%) for detecting *L pneumophila* serogroup 1, which is the most common cause of sporadic CAP and CAP due to foreign travel in the UK.[97]

**Debate: Patient groups for legionella testing**

The British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE) guidelines differ in their recommendations regarding the patient groups that should be tested for *legionella pneumonia* using urine antigen.#

- The BTS recommends testing only patients with high-severity CAP, patients with risk factors, and all patients with CAP during outbreaks,[1] whereas NICE recommends that clinicians consider testing in people with moderate- or high-severity CAP.[64]
- A later comparison by the BTS of the key recommendations in the two guidelines (BTS published in 2009 and NICE in 2014) concluded that there were no major differences between them and, where there were differences, clinicians should follow the NICE guideline instead of the BTS guideline.[96]
- In the comparison, the BTS also stated that its recommendation to test for legionella in patients with risk factors and all patients with CAP during outbreaks remains valid as this was not examined by NICE.[96]

If the *legionella* urine antigen test is positive, remember to order sputum cultures from respiratory samples (e.g., obtained from bronchoscopy) for *Legionella* species. This is to aid outbreak and source investigation to prevent further cases.[1]

**Polymerase chain reaction (PCR) and serological tests**

Use PCR of sputum or other respiratory tract samples for respiratory viruses (influenza A and B, parainfluenza 1-3, adenovirus, respiratory syncytial virus) and atypical pathogens (*Mycoplasma pneumoniae*, *Chlamydia pneumoniae*, *Chlamydia psittaci*, *Coxiella burnetii*, and *Pneumocystis jirovecii* [if at risk]) in patients with high-severity CAP:[1]

- If unresponsive to beta-lactam antibiotics
- If there is a strong suspicion of an ‘atypical’ pathogen.

Consider urine antigen, PCR of upper (e.g., nose and throat swabs) or lower (e.g., sputum) respiratory tract samples, or serological investigations in patients with moderate- or low-severity CAP:[1]

- During outbreaks (e.g., Legionnaires’ disease)
- During mycoplasma epidemics, or
Community-acquired pneumonia (non COVID-19)

### Diagnosis

- When there is a particular clinical or epidemiological reason
- If available, PCR is preferred over serological investigations.

#### In the community

**Do not order microbiological tests routinely** in patients presenting with CAP in the community as:[1] [64]

- These patients are not usually severely ill and are at low risk of death[1]
- Delays in transport of specimens to laboratory reduces the yield of bacterial pathogens (especially *S pneumoniae*) from sputum cultures, and results are often received too late by the general practitioner to have any impact on initial management.[1]

**Only consider ordering microbiological tests in the community if:**[1]

- The patient’s symptoms do not improve with empirical antibiotic therapy
  - Consider sputum examination
  - The patient has a persistent productive cough, especially if they also have malaise, weight loss, or night sweats, or risk factors for tuberculosis (e.g., ethnic origin, social deprivation, older patients, previous history of tuberculosis, contact history of tuberculosis)
  - Consider sputum examination for *Mycobacterium tuberculosis*
  - There is a clinical or epidemiological reason, such as an outbreak (e.g., Legionnaires’ disease) or during mycoplasma epidemics
  - Consider urine antigen, PCR of upper (e.g., nose and throat swabs) or lower (e.g., sputum) respiratory tract samples or serological investigations. If available, PCR is preferred over serological investigations.[1]

**Summary of the recommendations for microbiological investigations from the British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE)**[1] [64]
<table>
<thead>
<tr>
<th>CAP severity and treatment site</th>
<th>Preferred microbiological tests</th>
</tr>
</thead>
</table>
| **High-severity (CURB-65 = 3-5; CRB-65 = 3-4): treat in hospital** | • Blood and sputum (or other respiratory sample) cultures (plus Gram stain if available)  
• **Legionella and pneumococcal urine antigen** if legionella or pneumococcal pneumonia is suspected  
  • If legionella urine antigen is positive: sputum or other respiratory sample for legionella culture  
  • If legionella urine antigen is positive: sputum or other respiratory sample for legionella culture  
  • PCR of sputum (or other respiratory tract sample) for respiratory viruses (influenza A and B, parainfluenza 1-3, adenovirus, respiratory syncytial virus) and **atypical pathogens** (Mycoplasma pneumoniae, Chlamydia pneumoniae, Chlamydia psittaci, Coxiella burnetii, Pneumocystis jirovecii if at risk) if unresponsive to beta-lactam antibiotics and/or with a strong suspicion of an 'atypical' pathogen |
| **Moderate-severity (CURB-65 = 2; CRB-65 = 1-2): treat in hospital** | • Blood and sputum cultures (consider Gram stain if available)  
• **Legionella and pneumococcal urine antigen** if legionella or pneumococcal pneumonia is suspected  
  • If legionella urine antigen is positive: sputum or other respiratory sample for legionella culture  
  • **During outbreaks (e.g., Legionnaires' disease) or mycoplasma epidemic**: consider urine antigen, PCR* of upper (e.g., nose and throat swabs) or lower (e.g., sputum) respiratory tract samples or serological investigations |
| **Low-severity (CURB-65 = 0-1; CRB-65 = 0): treat at home or in hospital** | • None routinely  
• **During outbreaks (e.g., Legionnaires' disease) or mycoplasma epidemics**: consider urine antigen, PCR* of upper (e.g., nose and throat swabs) or lower (e.g., sputum) respiratory tract samples or serological investigations |

*If available, PCR is preferred over serological investigations.[1]
Practical tip

In routine clinical practice, pathogens are identified only in about one third to one quarter of patients with CAP admitted to hospital. Despite this, identifying the causative organism of CAP and sensitivity patterns is important because it:

- Allows for appropriate selection of antibiotic regimens. Change to targeted and narrow-spectrum antibiotic therapy is recommended once the pathogen is identified unless there are concerns of dual infection.
- Detects certain pathogens with public health significance and/or those that cause serious conditions that require different treatment from standard empiric antibiotics. These include:
  - *Legionella* species
  - Influenza A and B, including avian influenza A H5N1 and avian influenza A H7N9
  - Middle East respiratory syndrome coronavirus (MERS-CoV) and severe acute respiratory syndrome coronavirus (SARS-CoV)
  - Community-associated methicillin-resistant *S aureus* (CA-MRSA)
  - Agents of bioterrorism
  - Other emerging pathogens
- Allows for monitoring of the spectrum of organisms that cause CAP over time. This is important to establish sensitivity patterns.

Failure to improve

In hospital

Discuss with a senior colleague any patient who does not improve as expected. 

Consider repeat chest radiograph, C-reactive protein (CRP), white cell count, and further specimens for microbiology in patients not progressing satisfactorily after 3 days of treatment.

- A failure of CRP to fall by 50% or more at day 4 is associated with an increased risk for 30-day mortality, need for mechanical ventilation and/or inotropic support, and complications.

Practical tip

**Pointers to clinical improvement**: 

- Resolution of fever for >24 hours
- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
- No concerns over gastrointestinal absorption.

Consider referral to a respiratory physician.
In patients with high-severity CAP who are not responding to beta-lactam antibiotics or for whom an atypical or viral pathogen is suspected, order PCR (or other antigen detection test) of sputum or other respiratory tract sample.[1]

In the community

Consider assessing oxygenation via pulse oximetry if available.[1]

- Oxygen saturation <94% is an adverse prognostic factor in CAP and also may be an indication for oxygen therapy.[76]
- Consider referring these patients to hospital urgently. [1]

During recovery

Do not request a repeat chest radiograph before discharge from hospital in patients who have recovered satisfactorily from CAP.[1]

Request a repeat chest radiograph during recovery after about 6 weeks for patients (regardless of whether they have been admitted to hospital):[1]

- With persisting symptoms or physical signs
- Who are at higher risk of underlying malignancy (especially people who smoke and those aged >50 years).

Practical tip

Resolution of radiographic changes occurs relatively slowly after CAP and lags behind clinical recovery.[1]

Consider bronchoscopy in patients with persisting signs, symptoms, and radiological abnormalities at around 6 weeks after completing treatment.[1]

- Consider chest and upper abdomen CT scan in patients with persistent signs or symptoms or with chest radiograph changes prior to bronchoscopy (e.g., if lung cancer is suspected).[98]

Case history

Case history #1

A 54-year-old smoker with multiple comorbidities (diabetes, hypertension, coronary artery disease) presents with a 2-day history of a productive cough with yellow sputum, chest tightness, and fever. Physical examination reveals a temperature of 38.3°C (101°F), BP of 150/95 mmHg, heart rate of 85 bpm, and a respiratory rate of 20 breaths per minute. His oxygen saturation is 95% at rest; lung sounds are distant but clear, with crackles at the left base. Chest x-ray reveals a left lower lobe infiltrate.

Other presentations

Pneumonia can occur at any age, but the incidence increases significantly in old age, and pneumonia is a leading cause of illness and death in older patients. The clinical manifestations of pneumonia in elderly persons are often less intense than those in younger patients.[2] Atypical pathogens such as *Mycoplasma pneumoniae*, *Chlamydophila pneumophila*, and respiratory viruses can present in a subacute fashion with gradual onset of fever, non-productive cough, constitutional symptoms, relatively
normal white blood cell count, and absent or diffuse findings on lung examination. Patients with severe pneumococcal or *Legionella pneumophila* pneumonia often progress rapidly to respiratory failure.

### Risk factors

#### Strong

**age >65 years**

Incidence increases significantly with age. Very advanced age has been associated with higher mortality from CAP.[36]

**residence in a healthcare setting**

Approximately 10% to 18% of all patients hospitalised for pneumonia are nursing home residents. Mortality in these patients may reach 55%. Patients in residential homes who develop pneumonia have traditionally been considered to have healthcare-associated pneumonia (HCAP) and not CAP. However, this definition has been criticised because it is not able to distinguish patients at risk for resistant pathogens, and each patient ought to be evaluated individually.

**COPD**

Associated with a 2- to 4-fold increased risk of CAP.[6] Data from one study conducted in patients with CAP compared the outcome of patients with and without COPD and found that the presence of COPD was an independent risk factor for mortality.[39]

**exposure to cigarette smoke**

Colonisation with pathogenic bacteria is frequent in smokers and presents an increased risk of lung infections, especially pneumococcal pneumonia.[40] One study of bacterial pneumonia found that HIV-infected smokers had >80% higher risk of developing pneumonia than those who had never smoked.[12] Another study showed that current smokers with pneumococcal CAP often develop severe sepsis and require hospitalisation at a younger age despite having fewer comorbid conditions than older patients.[42] Passive smoking at home is a risk factor for CAP in people aged 65 years or older.[43]

**alcohol abuse**

There is clear evidence that alcohol consumption increases the risk for CAP. A meta-analysis of 14 studies found that people who consumed alcohol at all or in higher amounts had an 83% higher risk of CAP compared to people who consumed no alcohol or lower amounts (relative risk of 1.83).[44] Consumption of 24 g, 60 g, and 120 g of pure alcohol daily has been shown to result in a relative risk for incident CAP of 1.12 (95% CI, 1.02-1.23), 1.33 (95% CI, 1.06-1.67), and 1.76 (95% CI, 1.13-2.77), respectively, relative to non-drinkers.[45]

**poor oral hygiene**

Oral and respiratory bacteria in dental plaques are shed into the saliva and can then be aspirated into the lower respiratory tract to cause infection. Aspiration pneumonia is one of the most serious problems in older patients. Low-quality evidence suggests that professional oral health care measures (e.g., brushing, swabbing, denture cleaning, mouth rinses) may reduce mortality due to pneumonia in nursing home residents compared to usual care.[46]
use of acid-reducing drugs

CAP is one of the most common adverse effects associated with use of proton-pump inhibitors.[47] This is thought to be due to a decrease in gastric acid secretion, which allows pathogens to colonise the upper respiratory tract more easily. Outpatient use of these drugs is associated with a 1.5-fold increased risk of CAP.[48] H2 antagonists may also be associated with an increased risk of CAP.[49]

contact with children

Regular contact with children is associated with an increased risk of CAP.[50] Two studies have reported that having children in the household increases the adjusted odds ratio from 1.00 for households with no children to 3.2,[51] or 3.41[52] for households with 3 or more children.

Weak diabetes mellitus

Associated with a moderate increase in the risk of CAP. The main reasons are the increased risk of aspiration, hyperglycaemia, decreased immunity and impaired lung function, and coexisting morbidity.

One study found that diabetes (type 1 and type 2) was a risk factor for pneumonia-linked hospitalisation. Another study[53] reported that pre-existing diabetes was associated with a higher risk of death after hospitalisation for CAP compared with patients hospitalised for non-infectious illnesses.[54] The risk of severe pneumococcal bacteraemia is also higher in diabetic patients.[55]

chronic renal disease

A significant risk factor for mortality in patients with CAP.[56] [57]

chronic liver disease

It is known that bacterial infections occur in 32% to 34% of hospitalised patients with cirrhosis, and approximately 15% of these infections are pneumonia (the third most common cause of infection in these patients).[58] One study reported that chronic liver disease is a risk factor for pulmonary complications in patients hospitalised with pneumococcal pneumonia.[59]

use of opioids

A case-control study found that prescribed opioids, especially those with immunosuppressive properties or higher doses, are associated with an increased risk of CAP in people with and without HIV infection.[60]

History & examination factors

Key diagnostic factors
cough with increasing sputum production (common)

Symptoms of a lower respiratory tract infection such as cough are frequently seen in people with CAP. [1] [64]

- Cough is one of the most common symptoms present in patients with CAP.[1] Cough is usually productive with mucopurulent sputum.
**Community-acquired pneumonia (non COVID-19)**

**Diagnosis**

- The presence of **mucopurulent sputum** is associated with **bacterial pneumonia**.\[1\]
- Scant or watery sputum is associated with an **atypical pathogen**.\[1\]
- Older patients may not present with **cough** and are more likely to have non-specific symptoms (e.g., confusion) and may be afebrile.\[1\]

**dyspnoea (common)**

Dyspnoea is frequently seen in people with CAP. [1] \[64\]

- Dyspnoea is one of the most useful predictive symptoms of CAP in the community (together with fever, tachypnoea, pleuritic chest pain, and new/focalising signs on physical examination of chest) when compared with the gold standard of radiological diagnosis of CAP.\[1\]

**pleuritic chest pain (common)**

Pleuritic chest pain is frequently reported in people with CAP, [1] \[64\] occurring in 30% of patients. [99]

- Pleuritic chest pain is one of the most useful predictive symptoms of CAP in the community (together with fever, dyspnoea/tachypnoea, and new/focalising signs on physical examination of chest) when compared with the gold standard of radiological diagnosis of CAP.\[1\]

**rigors or night sweats (common)**

Rigors or night sweats are usually present in people with CAP, but are less common in older patients. [1] \[64\] #

**fever (common)**

Fever is commonly seen in people with CAP, although older people may be afebrile. [1] \[64\]

- A fever (>38°C (>100°F)) is one of the most useful predictive symptoms of CAP in the community (together with dyspnoea/tachypnoea, pleuritic chest pain, and new/focalising signs on physical examination of chest) when compared with the gold standard of radiological diagnosis of CAP.\[1\]
- Older people may be afebrile. \[1\]

**abnormal auscultatory findings (common)**

New focal chest signs are frequently present on examination in people with CAP. [1] \[64\]

- You may hear crackles, decreased breath sounds, dullness to percussion, and wheeze.
- Tachypnoea is one of the most useful predictive symptoms of CAP in the community (together with fever, dyspnoea, pleuritic chest pain, and new/focalising signs on physical examination of chest) when compared with the gold standard of radiological diagnosis of CAP.\[1\]

**confusion (common)**

Confusion is frequently seen in older people presenting with CAP. [1] \[64\]

- Older people with CAP often present with non-specific symptoms such as confusion or worsening of underlying diseases, and may be afebrile. [1] \[64\] \[65\]
• **Atypical presentations** (without obvious chest signs) of CAP may include confusion, such as in the case of *legionella pneumonia*, which may present as constitutional upset, diarrhoea, and confusion.[73]

**Risk factors (common)**

*Your history should cover the following risk factors to help assess the likelihood of CAP.*

[1] [64]

(*denotes a strong risk factor for CAP)

• **Age ≥65 years***
  
  • Incidence of CAP increases significantly with age. Advanced age is associated with a higher mortality from CAP.[10]

• **Residence in a nursing home***
  
  • Mortality rates due to pneumonia in nursing home residents have been reported to reach 55%. [78] [79]
  
  • Nursing home residents also have an increased risk of aspiration pneumonia.[80]

• **Contact with children***
  
  • Regular contact with children is associated with an increased risk of CAP.[50]

• **Respiratory chronic diseases**
  
  • **COPD***, *asthma, and bronchitis* are associated with a 2-fold to 4-fold increased risk of CAP.[6]
  
  • COPD is an independent risk factor for mortality in patients with CAP.[39]

• **Other chronic comorbidities**
  
  • **Chronic heart disease.** [6] [39]
  
  • **Diabetes** [6] [39] – the risk of severe pneumococcal bacteraemia is higher in people with diabetes.[55]

• **Alcohol use/misuse***
  
  • People who consume alcohol at all or in higher amounts have an 83% higher risk of CAP compared with people who consume no alcohol or lower amounts (relative risk of 1.83).[44] For every 10-20 g higher alcohol intake per day, there is an 8% increase in the risk of CAP.[44]

• **Smoking***
  
  • Smoking is an independent risk factor for developing CAP.[81]
  
  • Passive smoking at home is also a risk factor for CAP in people aged 65 years or older.[43] [81]

• **Poor oral hygiene**
Community-acquired pneumonia (non COVID-19)

**Diagnosis**

- Poor oral hygiene (particularly dental dysaesthesia and wearing dental prosthesis) may contribute to a higher risk of CAP in adults.[82]

  - **Proton pump inhibitors**
    - Associated with the occurrence of CAP.[47]

  - **H2 antagonists**
    - Associated with the occurrence of CAP.[49]

  - **Prescribed opioids**
    - In particular, immunosuppressive opioids are associated with CAP.[60]

**Other diagnostic factors**

- **myalgia (common)**
  
  Non-specific symptoms such as myalgia have been reported by people with CAP. [1]

  - Older people with CAP frequently present with non-specific symptoms and worsening of pre-existing conditions.[1]

- **malaise (common)**
  
  Non-specific symptoms such as malaise have been reported by people with CAP. [1]

  - Older people with CAP frequently present with non-specific symptoms and worsening of pre-existing conditions.[1]

- **anorexia (common)**
  
  Non-specific symptoms such as anorexia have been reported by people with CAP. [1]

  - Older people with CAP frequently present with non-specific symptoms and worsening of pre-existing conditions.[1]

- **lethargy (common)**
  
  Non-specific symptoms such as lethargy have been reported by people with CAP. [1]

  - Older people with CAP frequently present with non-specific symptoms and worsening of pre-existing conditions.[1]

- **worsening of pre-existing conditions (common)**
  
  - Older people frequently present with non-specific symptoms and worsening of pre-existing conditions.[1]

- **sore throat (uncommon)**
  
  Atypical presentations (without obvious chest signs) of CAP may include sore throat.
Community-acquired pneumonia (non COVID-19)

Diagnosis

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.\[73\]

**headache (uncommon)**

**Atypical presentations** (without obvious chest signs) of CAP may include headache.

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.\[73\]

**nausea (uncommon)**

**Atypical presentations** (without obvious chest signs) of CAP may include nausea.

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.\[73\]

**abdominal pain (uncommon)**

**Atypical presentations** (without obvious chest signs) of CAP may include abdominal pain.

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.\[73\]

**diarrhoea (uncommon)**

**Atypical presentations** (without obvious chest signs) of CAP may include diarrhoea.

- Mycoplasma pneumonia in young adults may present as sore throat, headache, nausea, abdominal pain, and diarrhoea.\[73\]
- Legionella pneumonia may present as constitutional upset, diarrhoea, and confusion.\[73\]
# Diagnostic tests

## 1st test to order

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>chest x-ray</td>
<td>new shadowing (consolidation)</td>
</tr>
</tbody>
</table>

A definitive diagnosis of CAP requires evidence of consolidation on chest x-ray. [1] [64] [75] Perform a chest x-ray in all patients presenting in hospital as soon as possible and within 4 hours of admission.

![Posterior-anterior chest radiograph showing right upper lobe consolidation in a patient with community-acquired pneumonia](image)


- If the chest x-ray shows atypical changes or complicated pneumonia (e.g., cavitation, pleural effusion, multifocal consolidation), consider other imaging investigations (see our section on Other tests to consider).

**Practical tip**

A high-quality chest radiograph is very important to ensure accurate diagnosis and to avoid inappropriate antibiotic prescribing. One study reported that 29% of hospitalised patients treated for CAP did not have radiographic abnormalities. [83]

Bear in mind that it is more difficult to obtain a high-resolution image from a person with class III obesity (BMI ≥40).

**Do not perform a chest x-ray in patients with suspected CAP seen in the community unless** [1]

- There is diagnostic doubt
- Progress following treatment is not satisfactory at review
### Test

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>• The patient is at risk of underlying lung pathology such as lung cancer.</strong></td>
<td></td>
</tr>
<tr>
<td><strong>pulse oximetry</strong></td>
<td>• may reveal low arterial oxygen saturation</td>
</tr>
<tr>
<td><em>Use pulse oximetry (preferably while breathing air) to assess oxygen saturation in hospital to inform supportive treatment.</em></td>
<td>• oxygen saturation &lt;94% in a patient with CAP is an adverse prognostic factor and may be an indication for oxygen therapy and urgent referral to hospital.[76] General practitioners should consider assessing oxygenation via pulse oximetry if available (e.g., if working in an out-of-hours setting).[1]</td>
</tr>
<tr>
<td><strong>arterial blood gas (ABG)</strong></td>
<td>may reveal low arterial oxygen saturation</td>
</tr>
<tr>
<td><em>Measure ABG in patients with CAP receiving oxygen therapy with an SpO(_2) &lt;94%, those with a risk of hypercapnic ventilatory failure (CO(_2) retention), and all patients with high-severity CAP.</em></td>
<td>• Patients may be hypoxaemic and require supplemental oxygen.</td>
</tr>
<tr>
<td></td>
<td>• Oxygen saturation &lt;94% in a patient with CAP is an adverse prognostic factor and may be an indication for oxygen therapy and urgent referral to hospital.[76]</td>
</tr>
<tr>
<td><strong>Practical tip</strong></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Always record the inspired oxygen concentration clearly as this is essential for interpreting blood gas results.</td>
</tr>
<tr>
<td><strong>urea and electrolytes</strong></td>
<td>• usually normal; elevated in patients with severe CAP</td>
</tr>
<tr>
<td><em>Request urea and electrolytes to inform disease severity and renal function in patients being investigated in hospital.</em>[1]</td>
<td>• <em>urea &gt;7 mmol/L (&gt;19.6 mg/dL) counts for 1 point in the 6-point CURB-65 score to assess severity</em>[1] [64]</td>
</tr>
<tr>
<td></td>
<td>• Chronic renal failure is a significant risk factor for mortality in patients with CAP.[57] General investigations are not necessary for the majority of patients with CAP who are managed in the community.[1] [64]</td>
</tr>
<tr>
<td><strong>full blood count</strong></td>
<td>• leukocytosis</td>
</tr>
<tr>
<td><em>Leukocytosis is often seen in people with CAP.</em></td>
<td>• <em>WBC count &gt; 15 x10(^9) /L indicates a bacterial aetiology (particularly pneumococcal,) although lower counts do not exclude a bacterial cause</em></td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>C-reactive protein (CRP)</strong></td>
<td>• elevated</td>
</tr>
</tbody>
</table>
### Test vs Result

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
</table>
| Order CRP as a baseline measurement and to help rule out other acute respiratory illnesses in patients being investigated in hospital. | • A failure of CRP to fall by 50% or more at day 4 is associated with an increased risk for 30-day mortality, need for mechanical ventilation and/or inotropic support, and complications.[91]  
  General investigations are not necessary for the majority of patients with CAP who are managed in the community.[1] [64] |
| • a level >100 mg/L makes pneumonia likely[90]                         | • a level <20 mg/L with symptoms for more than 24 hours makes the presence of pneumonia highly unlikely[90] |
| liver function tests                                                  | usually normal; abnormal in patients with underlying liver disease or legionella infection[1] |
| Take blood for a baseline measurement. Provides information about liver function. | • Chronic liver disease is a risk factor for pulmonary complications in patients hospitalised due to pneumococcal pneumonia.[59]  
  General investigations are not necessary for the majority of patients with CAP who are managed in the community.[1] [64] |
## Other tests to consider

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>blood culture</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>Order blood cultures, ideally before antibiotics are given, in all patients with moderate- or high-severity CAP (as determined by the CURB-65 score) presenting in hospital. [1] [64]</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>• Blood cultures can be highly specific in determining the microbial aetiology. [1] [64]</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>• Bacteraemia is also a marker of illness severity. However, many patients with CAP do not have associated bacteraemia. [1] Microbial causes of CAP that can be associated with bacteraemia include: [1]</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>• <em>Streptococcus pneumoniae</em> (identified in approximately 60% of positive blood cultures) [92]</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>• <em>Haemophilus influenzae</em> (identified in 2% to 13% of positive cultures) [92]</td>
<td>growth of causative bacterial pathogen</td>
</tr>
<tr>
<td>• <em>Staphylococcus aureus</em> and <em>Klebsiella pneumoniae</em>.</td>
<td>growth of causative bacterial pathogen</td>
</tr>
</tbody>
</table>

Do not order blood cultures in patients with confirmed CAP who have low-severity disease and no comorbid conditions. [1] Empirical antibiotic therapy is associated with a good prognosis in these patients. [1]

### Debate: Blood cultures#

There is debate around the practicality of ordering routine blood cultures in patients hospitalised with CAP. This is mainly due to low sensitivity, cost, and the fact that results hardly ever influence antimicrobial management.

- In a study of 355 patients admitted to hospital for CAP, the proportion of false-positive blood cultures was 10%, and the proportion of true positives was 9% (95% CI, 3.3% to 5.5%). [93]
- Antibiotic therapy was changed on the basis of blood culture results in only 5% of patients (95% CI, 3% to 8%). [93]
- However, despite these limitations, most experts still recommend blood cultures in patients with high-severity CAP. [1]

### sputum culture (± Gram stain)

Take sputum samples for culture (± Gram stain) in all patients with moderate- and high-severity CAP (as determined by the CURB-65 score) presenting in hospital before starting antibiotics, and in patients who do not improve regardless of disease severity. [1] [64]

Order Gram stain of sputum cultures in patients with high-severity CAP or complications if available in your local laboratory. Gram stain is an immediate indicator of the likely pathogen and can help with interpreting culture results. [1]
**Test**

<table>
<thead>
<tr>
<th>Do not order microbiological tests routinely in patients presenting with CAP in the community. [1] [64]</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Evidence: Sputum Gram stain</th>
</tr>
</thead>
</table>

A prospective study of 1390 patients with bacteraemic CAP found a sensitivity for sputum Gram stain of: [94]

- 82% for pneumococcal pneumonia
- 79% for *Haemophilus influenzae* pneumonia
- 76% for staphylococcal pneumonia.

Specificities ranged from 93% to 96%.

<table>
<thead>
<tr>
<th>Practical tip</th>
</tr>
</thead>
</table>

Carrying out routine sputum Gram stains for all patients is unnecessary. [1] The test has a low sensitivity and specificity, and often does not contribute to initial management. Problems include: [1]

- Patients may not be able to produce good specimens
- Prior exposure to antibiotics
- Delays in transport and processing of samples, which reduces the yield of bacterial pathogens
- Difficulty interpreting the results due to contamination of the sample by upper respiratory tract flora, which may include potential pathogens such as *Streptococcus pneumoniae* and ‘coliforms’ (especially in patients already given antibiotics).

<table>
<thead>
<tr>
<th>urinary antigen testing for legionella and pneumococcus</th>
</tr>
</thead>
</table>

Consider pneumococcal and legionella urine antigen tests in people with moderate- or high-severity CAP. [64] #Order legionella urine antigen testing in all patients with specific risk factors and during an outbreak (e.g, Legionnaires’ disease) or during epidemic mycoplasma years. [1]

- **Pneumococcal urinary antigen testing** is useful for diagnosing pneumococcal pneumonia in adults and is less affected than blood/sputum cultures by prior antibiotic therapy. [1]
- **Legionella urinary antigen** testing allows for rapid results early in the illness. [1]

- For all patients who are positive for legionella urine antigen, order sputum cultures from respiratory samples (e.g., obtained from bronchoscopy) for *Legionella* species. This is to aid outbreak and source investigation with the aim of preventing further cases. [1]
- Legionella antigen testing by enzyme immunoassay is highly specific (>95%) and sensitive (80%) for detecting *Legionella pneumophila* serogroup 1, which is the most...
<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td>Do not order microbiological tests routinely in patients presenting</td>
<td>common cause of sporadic CAP and CAP due to foreign travel in the UK.</td>
</tr>
<tr>
<td>with CAP in the community.</td>
<td>[97]</td>
</tr>
</tbody>
</table>

**Evidence: Urine antigen testing for pneumococcal pneumonia**

*Studies have shown significantly greater sensitivity rates for the pneumococcal urine antigen test than for routine blood or sputum cultures.* [95]

- Results remain positive in 80% to 90% of patients for up to 7 days after starting antimicrobial treatment. [95]

**polymerase chain reaction (PCR) and/or serological tests**

*Allows for rapid identification of the pathogen. Order PCR of sputum or other respiratory samples (e.g., nose and throat swabs) in patients with high-severity CAP:* [1]

- If unresponsive to beta-lactam antibiotics
- If there is a strong suspicion of a respiratory virus (i.e., influenza A and B, parainfluenza 1-3, adenovirus, respiratory syncytial virus) or an 'atypical' pathogen:
  - *Mycoplasma pneumoniae*
  - *Chlamydia pneumoniae*
  - *Chlamydia psittaci*
  - *Coxiella burnetii*
  - *Pneumocystis jirovecii* (if at risk).

**Consider PCR in patients with low- or moderate-severity CAP during outbreaks (e.g., Legionnaires’ disease) or during epidemic mycoplasma, or when there is a particular clinical or epidemiological reason.* [1]

**Do not order microbiological tests routinely in patients presenting with CAP in the community.* [1] [64]

**CT scan of chest**

*Consider a CT scan of the chest when there is diagnostic doubt: for example, if the chest x-ray is of poor quality or there is an ill-defined consolidation.* [1]

- Findings on chest x-ray that should prompt you to perform a CT scan include:
  - **Cavitation** – CT helps identify alternative diagnoses such as tuberculosis, lung cancer, pulmonary infarct, septic pulmonary emboli, infected bulla, lung abscess

- detection of viral/atypical pathogen antigens or antibodies

- may show cavitations, pleural effusion, multifocal consolidation, neoplasm
Community-acquired pneumonia (non COVID-19)

Diagnosis

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
</table>
| ![Chest radiograph showing left upper lobe cavitating pneumonia](image) | **Consolidation pattern (multifocal)** — CT helps identify alternative diagnoses such as staphylococcal infection, tuberculosis, aspiration pneumonia, allergic bronchopulmonary aspergillosis, cryptogenic organising pneumonia, or drug hypersensitivity reaction  
**Pleural effusion** — CT (in conjunction with chest ultrasound and guided aspiration) helps identify parapneumonic effusion, empyema, tuberculosis, lung cancer  
• Approximately 40% of patients who are hospitalised for pneumonia develop a parapneumonic effusion.[85] |

*From the collection of Dr Jonathan Bennett. Used with permission.*
<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Left-sided pleural effusion</strong></td>
<td><img src="image1" alt="Left-sided pleural effusion" /></td>
</tr>
<tr>
<td>From the collection of Dr R Light. Used with permission</td>
<td></td>
</tr>
<tr>
<td><strong>Diagnosis</strong></td>
<td></td>
</tr>
<tr>
<td>Consider a chest and upper abdomen CT scan in patients with</td>
<td></td>
</tr>
<tr>
<td>persistent signs and symptoms or changes on chest x-ray prior to</td>
<td></td>
</tr>
<tr>
<td>bronchoscopy to rule out alternative diagnoses.</td>
<td></td>
</tr>
<tr>
<td><strong>chest ultrasound</strong></td>
<td></td>
</tr>
<tr>
<td>Consider chest ultrasound following chest x-ray when there is</td>
<td>consolidation may be seen; parapneumonic effusion may be seen</td>
</tr>
<tr>
<td>diagnostic doubt. [1]</td>
<td></td>
</tr>
<tr>
<td>• Pleural effusion seen on chest x-ray may prompt you to perform a</td>
<td></td>
</tr>
<tr>
<td>chest ultrasound (with or without guided aspiration and chest x-ray)</td>
<td></td>
</tr>
<tr>
<td>to help make an alternative diagnosis such as tuberculosis, lung</td>
<td></td>
</tr>
<tr>
<td>cancer, or pulmonary embolism.</td>
<td></td>
</tr>
<tr>
<td><strong>Debate: Ultrasound in the diagnosis of CAP</strong></td>
<td></td>
</tr>
<tr>
<td>Although a chest radiograph showing new shadowing that cannot be</td>
<td></td>
</tr>
<tr>
<td>attributed to any other cause is the ‘gold-standard’ for the</td>
<td></td>
</tr>
<tr>
<td>diagnosis of pneumonia, it may not always be feasible in a</td>
<td></td>
</tr>
<tr>
<td>community setting and it involves exposure to radiation.</td>
<td></td>
</tr>
</tbody>
</table>
Community-acquired pneumonia (non COVID-19)

### Diagnosis

<table>
<thead>
<tr>
<th>Test</th>
<th>Result</th>
</tr>
</thead>
</table>
| • Emerging evidence has shown that lung ultrasound is a possible accurate diagnostic test for people with CAP. However, the benefits of its use in practice over chest radiography are still unclear.  
• A meta-analysis of 12 studies looking at the diagnostic accuracy of lung ultrasound in people with CAP found a sensitivity and specificity of 0.88 and 0.86, respectively.[88] However, there were limitations, such as the large variability in the findings and the lack of heterogeneity of the studies reviewed.  
• Further evidence is required before recommendations can be made. | |
| thoracocentesis and pleural fluid culture  
Consider thoracocentesis in all patients with pleural effusion as this can reveal an infected pleural space consistent with a parapneumonic effusion or empyema. [1]  
• A positive Gram stain of pleural fluid indicates an empyema. In these patients, drain pleural fluid in those with an empyema or clear pleural fluid with pH <7.2.[1] | exudate; growth of causative bacterial species in case of empyema |
| computer tomographic pulmonary angiography (CTPA)  
Consider CTPA to rule out pulmonary embolism if symptoms came on quickly (within minutes) or pain and breathlessness preceded infective symptoms. [87]  
• CTPA has the best diagnostic accuracy of all advanced non-invasive imaging methods in the detection of pulmonary embolism.[100] | may reveal a thrombus in a pulmonary artery; appears as a partial or complete intraluminal filling defect |
| bronchoscopy  
Consider bronchoscopy during recovery in patients with persisting signs and symptoms of CAP and radiological abnormalities at around 6 weeks after completing treatment. [1]  
• The most common techniques are bronchoalveolar lavage (BAL) and protected specimen brushing (PSB). | • BAL : $10^4$ colony-forming units (CFU)/mL indicates infection  
• PSB : $10^3$ CFU/mL has been recommended to distinguish colonisation from infection |
### Differential diagnosis

<table>
<thead>
<tr>
<th>Condition</th>
<th>Differentiating signs / symptoms</th>
<th>Differentiating tests</th>
</tr>
</thead>
</table>
| Coronavirus disease 2019 (COVID-19)            | • Residence in/travel to a country/area or territory with local transmission, or close contact with a confirmed or probable case of COVID-19, in the 14 days prior to symptom onset.  
• Signs and symptoms of viral pneumonia caused by COVID-19 and pneumonia caused by bacteria (either primary or secondary to COVID-19) are similar so it may be difficult to differentiate between the conditions clinically.  
• COVID-19 viral pneumonia may be more likely if the patient presents with a history of typical COVID-19 symptoms for about a week, severe myalgia, anosmia, breathlessness, and absence of pleuritic pain.  
• A bacterial cause of pneumonia may be more likely if the patient becomes rapidly unwell after only a few days of symptoms and presents with pleuritic pain, purulent sputum, and no history of typical COVID-19 symptoms.  
• This topic covers pneumonia caused by COVID-19 as a differential diagnosis only. For more detail on the diagnosis and management of community-acquired pneumonia caused by COVID-19, see our topic Coronavirus disease 2019 (COVID-19). | • Real-time reverse transcription polymerase chain reaction (RT-PCR): positive for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) RNA.  
• It is not possible to differentiate COVID-19 from other causes of pneumonia on chest imaging.                                                                                               |
<p>| Acute bronchitis                               | • No dyspnoea, no lung crackles, mild presentation. Often related to a viral upper respiratory tract infection.                                                                                                                                                                             | • No consolidation on CXR, with frequency related to viral infection.                                                                                                                                                       |</p>
<table>
<thead>
<tr>
<th>Condition</th>
<th>Differentiating signs / symptoms</th>
<th>Differentiating tests</th>
</tr>
</thead>
<tbody>
<tr>
<td>Congestive heart failure</td>
<td>• Peripheral oedema, cardiomegaly, hypotension.</td>
<td>• Bilateral interstitial pattern or pleural effusions seen on CXR.</td>
</tr>
<tr>
<td>COPD exacerbation</td>
<td>• Increased expectoration and cough, and worsening of dyspnoea against a background of COPD. Patient is often a smoker.</td>
<td>• CXR shows hyperinflation.</td>
</tr>
<tr>
<td>Asthma exacerbation</td>
<td>• Symptoms and signs of bronchospasm, with worsening of underlying lung disease.</td>
<td>• No consolidation on CXR.</td>
</tr>
<tr>
<td>Bronchiectasis exacerbation</td>
<td>• Increased expectoration and cough, and worsening of dyspnoea, with worsening of underlying lung disease. Infections are typically recurrent.</td>
<td>• No consolidation on CXR.</td>
</tr>
<tr>
<td>Tuberculosis</td>
<td>• Typically a long history, often with constitutional symptoms. Many patients will have lived in an endemic area.</td>
<td>• Cavitation on CXR, enlarged lymph nodes, positive purified protein derivative (PPD) skin testing.</td>
</tr>
<tr>
<td>Lung cancer or lung metastases</td>
<td>• Constitutional symptoms are common.</td>
<td>• Consolidation on CXR may be multiple, with pleural effusion commonly seen.</td>
</tr>
<tr>
<td>Empyema</td>
<td>• Constitutional symptoms are common, usually associated with a recent respiratory infection.</td>
<td>• Pleural effusion seen on CXR. Microbiology of pleural fluid may reveal infecting organism.</td>
</tr>
</tbody>
</table>
| Pulmonary embolism              | • Suspect pulmonary embolism in a patient with acute onset of dyspnoea, pleuritic chest pain, or features of deep vein thrombosis. In general, symptoms developing within minutes are more suggestive of pulmonary embolism than of community-acquired pneumonia.  
  • Cough is usually non-productive.  
  • Fever is generally lower in pulmonary embolism (i.e., below 39°C [102.2°F]).[73] | • Multiple-detector computed tomographic pulmonary angiography (CTPA): direct visualisation of thrombus in a pulmonary artery; appears as a partial or complete intraluminal filling defect. |
## Condition | Differentiating signs / symptoms | Differentiating tests
--- | --- | ---
**Pneumothorax** | • May be difficult to differentiate on the basis of signs and symptoms. In general, symptoms developing within minutes are more suggestive of pneumothorax than of community-acquired pneumonia.  
• Spontaneous pneumothorax may occur as a complication of pneumonia. | • CXR: presence of a visceral pleural line.[102]

**Hypersensitivity pneumonitis** | • May be difficult to differentiate on the basis of signs and symptoms.  
• Acute hypersensitivity pneumonitis lasts only a few days and recurs with each additional exposure. | • Immunological response to causative antigen: positive.

### Diagnostic criteria

Determine disease severity (and therefore mortality risk) in patients with a working diagnosis of pneumonia using the CURB-65 score in hospital or the CRB-65 score in the community together with your clinical judgement. The score allows initiation of appropriate antibiotic therapy and confirms whether the patient can be managed in the community or needs to be admitted to hospital.

#### CURB-65 score[103]

Recommended by the British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE) in the UK for use in the hospital setting,[1][64] CURB-65 stratifies patients according to the presence or absence of five prognostic features. Mortality at 30 days increases with the number of criteria that are met. Always use the CURB-65 score in conjunction with your clinical judgement.[1][64]

**Scoring of the CURB-65 for CAP in hospital**

- **Prognostic factors**
  - Confusion (e.g., Abbreviated Mental Test score ≤8 [Abbreviated Mental Test Score]): 1 point
  - Urea >7 mmol/L (>19.6 mg/dL): 1 point
  - Respiratory rate ≥30 breaths/minute: 1 point
  - Blood pressure <90 mmHg systolic or <60 mmHg diastolic: 1 point
  - Age ≥65 years: 1 point

- **Score**
  - **Score 3-5: high-risk; 30-day mortality >15%**
• Score of 3 or more: discuss with senior colleague at the earliest opportunity and manage as high-severity pneumonia.
• Score of 4-5: arrange emergency assessment by a critical care specialist.
• **Score 2: moderate-risk; 30-day mortality 3% to 15%**
  • Consider for short-stay inpatient treatment or hospital-supervised outpatient treatment.
• **Score 0-1: low-risk; 30-day mortality <3%**
  • Consider for outpatient treatment.

**CRB-65 score**[103]

Recommended by the BTS and NICE in the UK to be used in the community setting,[1] [64] CRB-65 stratifies patients according to the presence or absence of four prognostic features. Always use the CRB-65 score in conjunction with your clinical judgement.[1] [64]

Scoring of the CRB-65 for CAP in the community [103]

- **Prognostic factors**
  - Confusion (e.g., Abbreviated Mental Test score ≤8 [Abbreviated Mental Test Score]): 1 point
  - Respiratory rate ≥30 breaths/minute: 1 point
  - Blood pressure <90 mmHg systolic or <60 mmHg diastolic: 1 point
  - Age ≥65 years: 1 point

- **Score**
  - **Score 3-4: high-risk; 30-day mortality >10%**
    • Admit to hospital immediately.
  - **Score 1-2: moderate-risk; 30-day mortality 1% to 10%**
    • Consider hospital referral and assessment (particularly in those with a score of 2).
  - **Score 0: low-risk; 30-day mortality <1%**
    • Consider for treatment at home.

**Pneumonia severity index (PSI)**[104]

The PSI score predicts the risk of 30-day mortality; patients with a high risk are managed in hospital, and those with the highest risk are managed in the intensive care unit. The PSI stratifies patients into 5 categories based on patient age, comorbidities, physical examination, and results of laboratory testing. The principal limitation is the high score accorded to variables such as age and comorbidities. In the UK, the BTS and NICE consider the simplicity of the calculation of the CURB-65 score to be an advantage over PSI.[1] [64]

Scoring of the PSI for CAP [104]

- **Demographics**
Community-acquired pneumonia (non COVID-19)

**Diagnosis**

- Male: points = age in years
- Female: points = age in years minus 10
- Nursing home resident: +10 points
- Liver disease: +20 points
- Neoplastic disease: +30 points
- Congestive heart failure: +10 points
- Cerebrovascular disease: +10 points
- Renal failure: +10 points

- **Physical examination findings**
  - Altered mental status: +20 points
  - Respiratory rate ≥30 breaths/minute: +20 points
  - Systolic blood pressure <90 mmHg: +20 points
  - Temperature <35°C (<95°F) or ≥40°C (≥104°F): +15 points
  - Pulse rate ≥125 beats/minute: +10 points

- **Laboratory and radiographic findings**
  - Arterial pH <7.35: +30 points
  - Urea ≥10.7 mmol/L (≥30 mg/dL): +20 points
  - Sodium <130 mmol/L (<130 mEq/L): +20 points
  - Glucose ≥13.9 mmol/L (≥250 mg/dL): +10 points
  - Haematocrit <30%: +10 points
  - PaO₂ <60 mmHg (<90% O₂ saturation): +10 points
  - Pleural effusion: +10 points

- **Score**
  - Risk class I: 0 to 50 points: outpatients; 0.1% mortality
  - Risk class II: 51 to 70 points: outpatients; 0.6% mortality
  - Risk class III: 71-90 points: short hospital stay for observation; 2.8% mortality
  - Risk class IV: 91-130 points: hospital admission; 8.2% mortality
  - Risk class V: >130 points: hospital admission; 29.2% mortality
**Recommendations**

**Urgent**

*Think* Could this be sepsis? * whenever an acutely unwell person presents with likely infection, even if their temperature is normal.* Follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.[66] [67] [68]

- Remember that sepsis represents the severe, life-threatening end of infection.[69]
- Pneumonia is one of the main sources of infection in sepsis.[70]
- It is important to make an early, accurate diagnosis of infection and use antibiotics appropriately.

**Urgent: in hospital**

Stratify patients with pneumonia confirmed by chest x-ray according to mortality risk and disease severity using the CURB-65 score and your clinical judgement. Use the score to help inform your management plan.

- Score of 3-5: high-severity.
  - Score of 3 or more: discuss with a senior colleague at the earliest opportunity and manage as high-severity pneumonia.[1] [64]
  - Score of 4 or 5: arrange emergency assessment by a critical care specialist.[1]
- Score of 2: manage as moderate-severity pneumonia.
- Score of 0 or 1: manage as low-severity pneumonia.

Give empirical antibiotics immediately to all patients with CAP confirmed by chest x-ray. Patients should receive antibiotics within 4 hours of presentation to hospital.[1] [105]

- High-severity CAP: give broad-spectrum intravenous antibiotics – refer to your local protocol.
- Moderate- and low-severity CAP: give oral antibiotics (or intravenous if oral therapy is contraindicated) – refer to your local protocol.

Give empirical antibiotics to patients with life-threatening disease based on a presumptive clinical diagnosis, then order an immediate chest x-ray to confirm the diagnosis.[1]

- Follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.[66] [67] [68]

Assess oxygen requirements. Prescribe oxygen if oxygen saturation <94% and maintain at target range.[1] In patients at risk of hypercapnia prescribe oxygen if oxygen saturation <88%. [1]

- Target oxygen saturation range of:
  - 94% to 96% in acutely ill patients who are not at risk of hypercapnia[72]
  - 88% to 92% in patients at risk of hypercapnia.[71]
Measure and record observations at least twice daily and more frequently (e.g., every hour) in those admitted to a critical care unit to inform your management plan.[1]

Review all patients with high-severity CAP (high risk of death) at least every 12 hours until improvement.[1] This should be done by a senior colleague and the medical team. [1]

Urgent: in the community

Stratify patients according to mortality risk and disease severity using the CRB-65 score (see our Diagnosis – recommendations section for more information) and your clinical judgement.[1]

- Score of 3 or more (high-severity): admit patient to hospital immediately.[1][64]
- Score of 1 or 2 (moderate-severity): refer to hospital. These patients are at increased risk of death, particularly those with a score of 2.[1][64]
- Score of 0 (low-severity): treat most patients at home.[1][64]

Consider treatment at home for patients with a CRB-65 score of 0 (low-severity), or a CRB-65 score of 1 or 2 (medium-severity) if they wish to be treated at home and they meet all of the following criteria:[1][64]

- They are able to take oral medication safely and reliably
- Their social circumstances make them suitable for treatment at home
- They do not have unstable comorbidities.

Take a cautious approach when deciding whether it is safe to treat any patient with moderate-severity CAP (CRB-65 score of 1 or 2) in the community.

Give empirical antibiotics prior to hospital transfer (usually by blue-light ambulance in the UK) to patients with suspected CAP considered to be life-threatening.[1] Follow your local protocol.

Consider giving antibiotics prior to hospital transfer (usually by blue-light ambulance in the UK) to patients with suspected CAP where there are likely to be delays of over 6 hours to hospital admission and treatment.[1] Follow your local protocol.

Key Recommendations

Risk assessment and management of CAP in the first 4 hours: hospital setting
**Hospital setting:** manage according to clinical judgement + CURB-65 score

0 - 1
Low-severity (risk of death <3%)

Other reasons for admission (i.e., unstable comorbidities, social circumstances, patient’s wishes)

**NO**
- Home
  - Oral amoxicillin preferred (or intravenous if oral not possible)

**YES**
- Hospital
  - Order microbiology testing
  - Dual antibiotic therapy (amoxicillin plus clarithromycin preferred)

2
Moderate-severity (risk of death 9%)

Hospital

Supportive care

3 - 5
High-severity (risk of death 15% to 40%)

Hospital

Supportive care

Order microbiology testing

Intravenous empirical broad-spectrum antibiotics: amoxicillin/clavulanate plus clarithromycin preferred (if Legionella strongly suspected, consider adding levofloxacin)

Urgent senior colleague review

Consider ICU transfer (if CURB-65 scores = 4 and 5)

*Risk assessment and management of CAP in the first 4 hours: hospital setting*


*Risk assessment and management of CAP in the first 4 hours: community setting*
Community-acquired pneumonia (non COVID-19)

**Treatment**

**Community setting:** manage according to clinical judgement + CRB-65 score

- **0** Low-severity
  - Likely suitable for home treatment*
  - Oral amoxicillin preferred plus rest, smoking cessation, fluids, and analgesics if pain

- **1 - 2** Moderate-severity
  - Consider hospital referral**
  - Do not give antibiotics prior to hospital referral

- **3 - 4** High-severity
  - Urgent hospital admission
  - Prior to transfer: Consider giving intravenous benzylpenicillin or oral amoxicillin if life-threatening or if a delay in hospital treatment of >6 hours is likely

*Risk assessment and management of CAP in the first 4 hours: community setting. *Consider social circumstances, unstable comorbidities, ability to take oral treatment safely and reliably, and patient’s wishes when deciding whether to treat patients at home. **The majority of patients with moderate-severity CAP should be referred to hospital. However, you should consider social circumstances, ability to take oral treatment safely, unstable comorbidities, and patient’s wishes when deciding on home treatment#


**Supportive care**

Give **intravenous fluids** to patients with **volume depletion**.[1]

Arrange for patients with CURB-65 scores of 4 and 5 and an indication for intensive care unit (ICU) admission to be **transferred to ICU** and managed by ICU specialists together with respiratory physicians. [1]

- Patients with respiratory failure despite appropriate oxygen therapy require urgent airway management and possible intubation.

Do not routinely give **non-invasive ventilation (NIV)** or **continuous positive airways pressure (CPAP) support** in patients with respiratory failure due to CAP.[1]

- If indicated, you should conduct a trial of non-invasive support only in a critical care area where immediate expertise is available to allow a rapid transition to invasive ventilation.[1]
Give simple analgesia (e.g., paracetamol) as appropriate (e.g., for pleuritic pain).[1]

Other treatments
Do not give corticosteroids routinely to patients with CAP of any severity.[1] [64]

- Discuss with a senior colleague patients with comorbidities for which corticosteroids are indicated.[64]

Failure to improve
Discuss with a senior colleague any patient who does not improve as expected.[1]

- Consider repeat chest x-ray, C-reactive protein, white cell count, and further specimens for microbiology in patients who are not progressing satisfactorily after 3 days of treatment.[1]
- Consider referral to a respiratory physician.[1]

Prognosis
For hospitalised patients, mortality rate ranges from 5% to 15%, but increases to between 20% and 50% in patients requiring admission to the ICU.[6] [106] Patients treated in the community generally have a good prognosis.[1]

Full Recommendations
Risk stratification
Determine disease severity in patients with a working diagnosis of CAP as severe pneumonia is associated with a high risk of complications and death. [65]

- Stratify patients into those with low-, moderate- or high-severity disease (see below).[1] [64]
- Early identification of severity allows initiation of appropriate antibiotic therapy, as well as confirming whether the patient can be managed in the community or needs to be admitted to hospital where assisted ventilation in an intensive care setting may be necessary.[1] [64]

Practical tip
Think ‘Could this be sepsis?’ whenever an acutely unwell person presents with likely infection, even if their temperature is normal. Follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigation and treatment of all patients with suspected sepsis, or those at risk, within 1 hour.[66] [67] [68]

- Remember that sepsis represents the severe, life-threatening end of infection.[69]
- Pneumonia is one of the main sources of infection in sepsis.[70]
- It is important to make an early, accurate diagnosis of infection and use antibiotics appropriately.

Use your clinical judgement in conjunction with a scoring system.[1] [64]

- CURB-65 in hospital
- CRB-65 in the community.
Debate: The importance of clinical judgement

Your clinical judgement is key when assessing severity in CAP using a scoring system.

• For example, consider a young person with a normal blood pressure and urea level, but a low oxygen saturation despite supplemental oxygen. This person could potentially have severe pneumonia, but you could miss this if you only use a scoring system in your assessment.[65]
• You should use mortality prediction tools such as CURB-65 and CRB-65 to supplement, rather than replace, clinical judgement when assessing severity in CAP.

Consult with a senior colleague regarding the decision to start antibiotics at the earliest opportunity. [1]

In hospital

Assess severity using the CURB-65 score and your clinical judgement to identify patients with suspected CAP at high risk of death so that you can prioritise immediate treatment and consider whether the patient should be admitted.[1] [64]

Assess severity regularly in all patients with CAP following hospital admission.[1]

• An early opportunity for this is by a senior colleague and the medical team during the ward round following admission.[1]

CURB-65 score: hospital setting#

Score 1 point for each feature present:

• Confusion*
• Urea >7 mmol/L (>19.6 mg/dL)
• Respiratory rate ≥30/minute
• Blood pressure (systolic <90 mmHg or diastolic ≤60 mmHg)
• Age ≥65 years.

• *Use the Abbreviated Mental Test to assess for confusion. [Abbreviated Mental Test Score]
  • Each question scores 1 mark, with a total possible score of 10 marks.
  • A score of 8 or less has been used to define mental confusion in the CURB-65 severity score.[64]

Discuss patients with a CURB-65 score of 3 or more with a senior colleague at the earliest opportunity and manage as high-severity pneumonia (see below).[1] [64]

Arrange emergency assessment of patients with a CURB-65 score of 4 or 5 by a critical care specialist.[1]

Risk assessment and management of CAP in the first 4 hours: hospital setting
Follow your local protocol when prescribing antibiotics. The recommendations given here are based on British Thoracic Society (BTS) guidelines.[1] See our Management in hospital section below for more detail on antibiotic regimens.

Evidence: Effectiveness of the CURB-65 mortality risk score

CURB-65 is an accurate prognostic tool for predicting mortality in patients with CAP, and it is recommended by the British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE) alongside clinical judgement. [1][64]

- It was developed from three prospective studies in the UK, New Zealand, and the Netherlands that included more than 1000 patients with CAP.[103]
- Validation studies have been carried out in several countries (including more than 3000 patients in total). They have shown increasing mortality with increasing CURB-65 scores, ranging...
from 0% to 1.1% (CURB-65 score = 0) to 17% to 60% (CURB-65 score = 5),[107][108][109][110] and increasing need for mechanical ventilation with increasing CURB-65 scores, ranging from 0% (CURB-65 score = 0) to 11% (CURB-65 score = 5).[110]

• In a study comparing CURB-65 to the pneumonia severity index (PSI), CURB-65 showed equal sensitivity and higher specificity (74.6% CURB-65 vs. 52.2% PSI) in predicting mortality due to CAP.[107]
• Another study suggests that PSI may be more accurate than CURB-65 (and CRB-65) in predicting 30-day mortality in patients with CAP.[111]
• Despite emerging evidence for some benefits of PSI, both NICE and the BTS consider the simplicity of the calculation of the CURB-65 score to be an advantage over PSI. [1][64]

In the community
Assess severity using the CRB-65 score together with your clinical judgement to decide whether to treat patients with suspected CAP at home or refer to hospital for assessment or hospital admission. [1][64]

CRB-65 score: community setting

Score 1 point for each feature present:

• Confusion*
• Respiratory rate ≥30/minute
• Blood pressure (systolic <90 mmHg or diastolic ≤60 mmHg)
• Age ≥65 years.

*Use the Abbreviated Mental Test to assess for confusion. [Abbreviated Mental Test Score]

• Each question scores 1 mark, with a total possible score of 10 marks.
• A score of 8 or less has been used to define mental confusion in the CRB-65 severity score.[64]

Risk assessment and management of CAP: community setting
Community setting: manage according to clinical judgement + CRB-65 score

**Community-acquired pneumonia (non COVID-19)**

**TREATMENT**

**Risk assessment and management of CAP in the first 4 hours: community setting.** Consider social circumstances, unstable comorbidities, ability to take oral treatment safely and reliably, and patient’s wishes when deciding whether to treat patients at home. **The majority of patients with moderate-severity CAP should be referred to hospital. However, you should consider social circumstances, ability to take oral treatment safely, unstable comorbidities, and patient’s wishes when deciding on home treatment.**


Follow your local protocol when prescribing antibiotics. The recommendations given here are based on BTS and NICE guidelines.[1] [105] See our Management in the community section below for more detail on antibiotic regimens.

**Management in hospital**

**Antibiotics**

Give empirical antibiotics to all patients with CAP confirmed by chest x-ray immediately after diagnosis and within 4 hours of presentation to hospital.[1] [105]

- Aim to give antibiotics to all patients with confirmed CAP before they leave the initial assessment area.[1]
- Give empirical antibiotics to patients with life-threatening disease based on a presumptive clinical diagnosis of CAP, then order an immediate chest x-ray to confirm the diagnosis.[1]
Community-acquired pneumonia (non COVID-19)

Treatment

- For these patients, follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigating and treating all patients with suspected sepsis, or those at risk, within 1 hour.[66] [67] [68]
- Pneumonia is one of the main sources of infection in sepsis.[70]
- It is important to make an early, accurate diagnosis of infection and use antibiotics appropriately.

Consult with a senior colleague about your decision to start antibiotics at the earliest opportunity.[1]

  - Clearly document the indication for prescribing antibiotics in the medical notes.[1]

De-escalate treatment as soon as appropriate, including switching from intravenous to oral antibiotic therapy.[1] When making this decision consider response to treatment (see practical tip), change in disease severity, and contraindications to oral administration such as:[1]

  - Patient is unable to swallow (e.g., impaired swallowing reflex, impaired consciousness)
  - Gastrointestinal malabsorption for functional or anatomical reasons.

Review route of administration initially on the ward round following admission and then daily thereafter.[1]

Practical tip

Pointers to clinical improvement
The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]

- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of fever for >24 hours
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
- No concerns over gastrointestinal absorption.

Consider narrowing the spectrum of antibiotics or switching to pathogen-targeted antibiotics once a causative pathogen is identified.[1]

Empirical antibiotics

Treat the majority of patients empirically as the causative pathogen is only rarely identified at the initial assessment.[1]

Prescribe an appropriate antibiotic regimen according to your local protocol to help reduce the development of antibiotic resistance and Clostridium difficile infection.[1] To aid antibiotic stewardship, the British Thoracic Society (BTS) recommends:[1]
• Give empirical broad-spectrum intravenous antibiotics only to patients with high-severity CAP, and de-escalate to narrow-spectrum antibiotics as soon as clinically appropriate, based on the results of early microbiological investigations.[1]

  • This group comprises approximately one third of all patients admitted to hospital with confirmed CAP.[1]
  
• Give empirical antibiotics to all other patients and switch to pathogen-targeted antibiotics as soon as specific pathogens are identified.

• Consult with a microbiologist about appropriate antibiotic therapy.[1]

Consult local antibiotic protocols to determine the most appropriate choice based on local pathogen prevalence and antibiotic resistance patterns.

Consider whether a correct diagnosis of CAP has been made if a patient does not respond to initial empirical antibiotics. [1]

• Consider clinical and radiographic review to look for secondary diagnoses or complications of CAP such as pleural effusion/empyema, lung abscess, or worsening pneumonic shadowing.[1]

• Consider changing initial empirical antibiotics but first consider compliance with and adequate absorption of an oral regimen.[1]

The following recommendations are based on guidelines from the BTS and the National Institute for Health and Care Excellence (NICE).[1] [105]

High-severity CAP (CURB-65 score: 3 or more)

Always manage patients with high-severity CAP in hospital. [1]

Give empirical broad-spectrum intravenous antibiotics immediately after diagnosis and within 4 hours of presentation to hospital. [1] [105]

Evidence: Time to first antibiotic within 4 hours of presentation

The BTS CAP care bundle study was carried out in 2013 in 16 UK hospital trusts.

• The CAP care bundle consisted of four elements:

  • A chest x-ray obtained within 4 hours of hospital admission in all adults with suspected CAP
  • Oxygen assessment and prescription in keeping with the BTS oxygen guideline[89]
  • Severity assessment supported by the CURB-65 score
  • Timely and targeted antibiotics given according to CAP severity within 4 hours of admission.

• The study compared patients with CAP receiving the CAP care bundle with those receiving standard care. Analysis of data from 2118 adults (median age 75.3 years) showed that significantly more patients in the CAP care bundle group received antibiotics within 4 hours of admission (adjusted odds ratio [OR] 1.52, 95% CI 1.08 to 2.14, P = 0.016) and that 30-day
inpatient mortality was lower in this group (8.8% vs. 13.6%; adjusted odds ratio [OR] 0.59, 95% CI 0.37 to 0.95, \( P = 0.03 \)).[71]

Give a broad-spectrum beta-lactamase-resistant penicillin (e.g., amoxicillin/clavulanate) plus a macrolide (e.g., clarithromycin).[1]

- The BTS guideline recommends adding a fluoroquinolone to the existing empirical regimen (i.e., triple therapy) if the patient does not respond, or if legionella pneumonia is strongly suspected.[1] However, in practice there are concerns about the risk of using a macrolide and fluoroquinolone together as they can both prolong the QT interval. Some clinicians therefore replace the macrolide in the original empirical regimen with a fluoroquinolone instead (i.e., dual therapy). Consider the safety issues associated with fluoroquinolone use. Consult with a microbiologist and senior colleague before treating these patients.

- More recent (2019) guidelines from NICE on antimicrobial prescribing in adults also recommend amoxicillin/clavulanate plus clarithromycin (or erythromycin) as first-line in people with high-severity CAP. These recommendations are based mainly on expert opinion as the evidence is limited.[105] For patients who are allergic to penicillin, give a second-generation cephalosporin (e.g., cefuroxime) or a third-generation cephalosporin (e.g., cefotaxime or ceftriaxone) plus a macrolide (e.g., clarithromycin).[1]

- A small number of patients are allergic to both penicillins and cephalosporins; consult an infectious disease consultant for selection of appropriate antibiotics in these patients.

- NICE guidelines on antimicrobial prescribing in adults recommend levofloxacin (after considering safety issues associated with fluoroquinolone use) as an alternative antibiotic for patients with high-severity CAP who are allergic to penicillin.[105]

Review the need for intravenous antibiotics initially during the ward round following admission and then every day thereafter.[1]

Switch to oral antibiotics as soon as clinical improvement occurs (see practical tip above for pointers to clinical improvement), and as long as there are no contraindications to oral administration such as:[1]

- Patient is unable to swallow (e.g., impaired swallowing reflex, impaired consciousness)
- Gastrointestinal malabsorption for functional or anatomical reasons.

Give antibiotic therapy for 5 days. NICE recommends: [105]

- Prescribing the shortest course that is likely to be effective to reduce the risk of antimicrobial resistance and minimise the risk of adverse effects[105]
- Stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable. This should be based on your clinical judgement and the following criteria:[105] [112]

- Fever in the past 48 hours, or more than one sign of clinical instability:
  - Systolic blood pressure <90 mmHg
  - Heart rate >100/minute
  - Respiratory rate >24/minute
Community-acquired pneumonia (non COVID-19)

TREATMENT

- Arterial oxygen saturation <90% or PaO$_2$ <60 mmHg in room air.

In some people, longer courses might be needed due to individual circumstances. In the UK, some hospitals require consultation with the microbiology team if considering extending the duration of antibiotic treatment beyond 5 days in high-severity CAP. Follow your local protocol.

- NICE guidelines do not provide an upper limit on antibiotic course length. This will be determined in individual circumstances, based on time taken to reach clinical stability.[105]

Moderate-severity CAP (CURB-65 score: 2)

Give antibiotics as soon as possible after diagnosis and within 4 hours of presentation to hospital. [1] [105]

Consider patients with moderate-severity CAP for short-stay inpatient treatment or hospital-supervised outpatient treatment.[1]

Evidence: Time to first antibiotic within 4 hours of presentation

The BTS CAP care bundle study was carried out in 2013 in 16 UK hospital trusts.

- The CAP care bundle consisted of four elements:
  - A chest x-ray obtained within 4 hours of hospital admission in all adults with suspected CAP.
  - Oxygen assessment and prescription in keeping with the BTS oxygen guideline[89]
  - Severity assessment supported by the CURB-65 score
  - Timely and targeted antibiotics given according to CAP severity within 4 hours of admission.

- The study compared patients with CAP receiving the CAP care bundle with those receiving standard care. Analysis of data from 2118 adults (median age 75.3 years) showed that significantly more patients in the CAP care bundle group received antibiotics ≤4 hours after admission (adjusted OR 1.52, 95% CI 1.08 to 2.14, P = 0.016) and that 30-day inpatient mortality was lower in this group (8.8% vs. 13.6%; adjusted OR 0.59, 95% CI 0.37 to 0.95, P = 0.03).[71]

Most patients with moderate-severity CAP can be treated with dual oral antibiotic therapy. The preferred choice is amoxicillin plus a macrolide (e.g., clarithromycin).[1]

- NICE guidelines on antimicrobial prescribing in adults recommend monotherapy with amoxicillin in people with moderate-severity CAP, with the addition of clarithromycin (or erythromycin) only if an atypical pathogen is suspected. These recommendations are based mainly on expert opinion as the evidence is limited.[105]

Consider monotherapy with a macrolide for patients who have been treated in the community and who have not responded to an adequate course of amoxicillin prior to hospital admission.[1]
Community-acquired pneumonia (non COVID-19)

Treatment

- Deciding whether the course of amoxicillin was adequate is tricky and involves clinical judgement. Consult a senior clinician before prescribing monotherapy within the first 24 hours of admission.[1]

If oral antibiotics are contraindicated (e.g., patient is unable to swallow or has gastrointestinal malabsorption for functional or anatomical reasons), give intravenous amoxicillin or benzylpenicillin, plus clarithromycin.[1]

For patients who are allergic to penicillin or macrolides, consider oral doxycycline.[1] Alternative choices include oral levofloxacin or moxifloxacin (after considering safety issues associated with fluoroquinolone use).[1]

- NICE guidelines on antimicrobial prescribing in adults with CAP recommend doxycycline or clarithromycin in people who are allergic to penicillin.[105]

For patients who are allergic to penicillin in whom oral antibiotics are contraindicated, give a second-generation cephalosporin (e.g., cefuroxime) or a third-generation cephalosporin (e.g., cefotaxime or ceftriaxone) plus clarithromycin, or intravenous levofloxacin as monotherapy.[1]

If the patient does not respond to a combination of amoxicillin plus clarithromycin consider changing treatment to doxycycline or a fluoroquinolone with effective pneumococcal cover.[1]

More info: EMA and MHRA restrictions on the use of fluoroquinolone antibiotics

In November 2018, the European Medicines Agency (EMA) completed a review of serious, disabling, and potentially irreversible adverse effects associated with systemic and inhaled fluoroquinolone antibiotics. These adverse effects include tendonitis, tendon rupture, arthralgia, neuropathies, and other musculoskeletal or nervous system effects.

- As a consequence of this review, the EMA now recommends that fluoroquinolone antibiotics be restricted for use in serious, life-threatening bacterial infections only. Furthermore, they recommend that fluoroquinolones should not be used for mild to moderate infections unless other appropriate antibiotics for the specific infection cannot be used, and should not be used in non-severe, non-bacterial, or self-limiting infections. Patients who are older, have renal impairment, or have had a solid organ transplant, and those being treated with a corticosteroid are at a higher risk of tendon damage. Co-administration of a fluoroquinolone and a corticosteroid should be avoided.[113] The UK-based Medicines and Healthcare products Regulatory Agency (MHRA) supports these recommendations.[114]

- For this reason, fluoroquinolones should only be considered in moderate-severity CAP when it is considered inappropriate to use other antibiotics that are commonly recommended for the treatment of CAP. Consult with a microbiologist about whether a fluoroquinolone is an appropriate option for your patient.

Review the need for intravenous antibiotics initially on the ward round following admission and then every day thereafter.[1]

- Switch to oral antibiotics as soon as clinical improvement occurs (see practical tip), and as long as there are no contraindications to oral administration such as:[1]

  - Patient is unable to swallow (e.g., impaired swallowing reflex, impaired consciousness)
Community-acquired pneumonia (non COVID-19)

Treatment

• Gastrointestinal malabsorption for functional or anatomical reasons.

Practical tip

Pointers to clinical improvement
The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]
• Pulse rate <100 beats/minute
• Resolution of tachypnoea
• Clinically hydrated and taking oral fluids
• Resolution of fever for >24 hours
• Resolution of hypotension
• Absence of hypoxia
• Improving white cell count
• Non-bacteraemic infection
• No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
• No concerns over gastrointestinal absorption.

Give antibiotic therapy for 5 days. NICE recommends:[105]
• Prescribing the shortest course that is likely to be effective to reduce the risk of antimicrobial resistance and minimise the risk of adverse effects
• Stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable. This should be based on your clinical judgement and the following criteria:[1][112]
• Fever in past 48 hours, or more than one sign of clinical instability:
  • Systolic blood pressure <90 mmHg
  • Heart rate >100/minute
  • Respiratory rate >24/minute
  • Arterial oxygen saturation <90% or PaO₂ <60 mmHg in room air.

In some people, longer courses might be needed due to individual circumstances. In the UK, some hospitals require consultation with the microbiology team if considering extending the duration of antibiotic treatment beyond 5 days in moderate-severity CAP. Follow your local protocol.

• NICE guidelines do not provide an upper limit on antibiotic course length. This will be determined in individual circumstances, based on time taken to reach clinical stability.[105]

Low-severity CAP (CURB-65 score: 0-1)
Give antibiotics as soon as possible and within 4 hours of presentation to hospital.[1][105]

Evidence: Time to first antibiotic within 4 hours of presentation

The BTS CAP care bundle study was carried out in 2013 in 16 UK hospital trusts.
Community-acquired pneumonia (non COVID-19)

### Treatment

- The CAP care bundle consisted of four elements:
  - A chest x-ray obtained within 4 hours of hospital admission in all adults with suspected CAP
  - Oxygen assessment and prescription in keeping with the BTS oxygen guideline[89]
  - Severity assessment supported by the CURB-65 score
  - Timely and targeted antibiotics given according to CAP severity within 4 hours of admission.

- The study compared patients with CAP receiving the CAP care bundle with those receiving standard care. Analysis of data from 2118 adults (median age 75.3 years) showed that significantly more patients in the CAP care bundle group received antibiotics ≤4 hours after admission (adjusted OR 1.52, 95% CI 1.08 to 2.14, P = 0.016) and that 30-day inpatient mortality was lower in this group (8.8% vs. 13.6%; adjusted OR 0.59, 95% CI 0.37 to 0.95, P = 0.03).[71]

Most patients with **low-severity CAP** who are managed in hospital can be treated with oral antibiotics.[1] The **preferred choice is amoxicillin**.[1] Consider a macrolide (e.g., clarithromycin) or a tetracycline (e.g., doxycycline) for patients who are allergic to penicillin.[1]

- NICE recommends clarithromycin (or erythromycin) or doxycycline as alternatives to amoxicillin for patients allergic to penicillin and for patients in whom amoxicillin is unsuitable (e.g., if atypical pneumonia is suspected). NICE recommendations are based mainly on expert opinion as the evidence is limited.[105]

If the oral route is contraindicated (e.g., impaired swallowing reflex, impaired consciousness, gastrointestinal malabsorption), consider **intravenous amoxicillin, benzylpenicillin, or clarithromycin**.[1]

- Review the need for intravenous antibiotics initially during the ward round following admission and then every day after.[1]
- **Switch to oral antibiotics as soon as clinical improvement occurs** (see practical tip), and as long as there are no contraindications to oral administration.[1][105]

#### Practical tip

**Pointers to clinical improvement**

The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]

- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of fever for >24 hours
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
If the patient does not respond to amoxicillin monotherapy, consider switching to, or adding, a macrolide (e.g., clarithromycin). [1]

Give antibiotic therapy for 5 days. NICE recommends [105]

- Prescribing the shortest course that is likely to be effective to reduce the risk of antimicrobial resistance and minimise the risk of adverse effects
- Stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable. This should be based on your clinical judgement and the following criteria: [105] [112]
  - Fever in past 48 hours, or more than one sign of clinical instability:
    - Systolic blood pressure <90 mmHg
    - Heart rate >100/minute
    - Arterial oxygen saturation <90% or PaO₂ <60 mmHg in room air.

Pathogen-targeted antibiotic treatment

Switch from empirical antibiotics to pathogen-targeted antibiotics as soon as specific pathogens are identified (unless there are legitimate concerns about dual-pathogen infection). [1]

- Consult with a microbiologist about appropriate antibiotic therapy. [1]
- Only about one third to one quarter of patients with CAP admitted to hospital will have their pneumonia defined microbiologically. [1] Among these patients:
  - Around 14% have an atypical pathogen, of which:[21]
    - 7% have *Mycoplasma pneumoniae*
    - 4% have *Chlamydophila pneumoniae*
    - 3% have *Legionella pneumophila*
  - Those with infections due to organisms such as *Mycoplasma*, *Chlamydophila*, and *Coxiella burnetii* will be diagnosed late in the illness on the basis of seroconversion, reducing the opportunity for early targeted therapy. [1] Among patients managed in the community, very few will be microbiologically defined. [1]

Consider switching treatment once the results of sensitivity testing are available or following consultation with a microbiologist, intensivist, or respiratory physician. [1]

BTS recommendations for pathogen-targeted antibiotics [1]
<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Mycoplasma pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Legionella</em> species</td>
<td>Fluoroquinolone (orally or intravenously)</td>
<td>Clarithromycin (orally or intravenously) or azithromycin (in countries where it is used for managing pneumonia)</td>
</tr>
<tr>
<td><em>Streptococcus pneumoniae</em></td>
<td>Amoxicillin (orally) or benzylpenicillin (intravenously)</td>
<td>Clarithromycin (orally) or cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia psittaci</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Coxiella burnetii</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Haemophilus influenzae</em></td>
<td>Non-beta-lactamase-producing : amoxicillin (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td></td>
<td>Beta-lactamase-producing : amoxicillin/clavulanate (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td>Gram-negative enteric bacilli</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
<td>Fluoroquinolone (intravenously) or imipenem/cilastatin (intravenously) or meropenem (intravenously)</td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em></td>
<td>Ceftazidime (intravenously)</td>
<td>Ciprofloxacin (intravenously) or piperacillin/tazobactam (intravenously)</td>
</tr>
<tr>
<td></td>
<td><strong>plus</strong></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Gentamicin or tobramycin (dose monitoring required)</td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em> : non-MRSA</td>
<td>Flucloxacin (intravenously)</td>
<td></td>
</tr>
<tr>
<td></td>
<td><strong>with or without</strong></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em> : MRSA</td>
<td>Vancomycin (intravenously; dose monitoring required)</td>
<td></td>
</tr>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
<td>Alternative antibiotic</td>
</tr>
<tr>
<td>----------</td>
<td>----------------------</td>
<td>------------------------</td>
</tr>
<tr>
<td></td>
<td>linezolid (intravenously) or teicoplanin (intravenously)</td>
<td>with or without Rifampicin (orally or intravenously)</td>
</tr>
</tbody>
</table>

**Supportive care**

Provide **supportive care for patients treated** in hospital. **This may include the following measures.** [1]

**Oxygen**

**Assess oxygen saturation in all patients by pulse oximetry (preferably while breathing air).**

- Give oxygen if oxygen saturation <94% and maintain at target range.[1] In patients at risk of CO<sub>2</sub> retention give oxygen if oxygen saturation <88%. Early oxygen assessment is associated with improved prognosis.[71]
- **BTS guidelines recommend a target oxygen saturation range of :**[71]
  - 94% to 98% in age >16 years
  - 88% to 92% in patients at risk of hypercapnia.

However, **latest evidence** suggests that liberal use of supplemental oxygen (target SpO<sub>2</sub> >96%) in acutely ill adults is associated with higher mortality than more conservative oxygen therapy.[72] A reasonable approach is to aim for a **target oxygen saturation of 94% to 96%** in acutely ill patients who are not at risk of hypercapnia.

**Evidence: Target oxygen saturation in acutely ill adults**

*Too much supplemental oxygen increases mortality.*

**Latest evidence supports a 96% upper limit for target oxygen saturation in non-hypercapnic acutely ill adults.**

- Guidelines differ in their recommendations on target oxygen saturation in acutely unwell adults who are receiving supplemental oxygen. The 2017 BTS guideline recommends a target SpO<sub>2</sub> range of 94% to 98% for patients not at risk of hypercapnia,[89] whereas the 2015 Thoracic Society of Australia and New Zealand guideline recommends 92% to 96%.[115]
- A 2018 systematic review including meta-analysis of data from 25 randomised controlled trials found that, in adults with acute illness, liberal oxygen therapy (broadly equivalent to a target saturation >96%) is associated with higher mortality than conservative oxygen therapy (broadly equivalent to a target saturation ≤96%).[72] In-hospital mortality was 11 per 1000 higher for the liberal oxygen therapy group versus the conservative therapy group (95% CI, 2-22 per 1000 more). Mortality at 30 days was also higher in the group
Community-acquired pneumonia (non COVID-19)

who had received liberal oxygen (RR 1.14, 95% CI 1.01 to 1.29). The trials included adults with sepsis, critical illness, stroke, trauma, myocardial infarction, or cardiac arrest, and patients who had had emergency surgery. The review excluded studies limited to people with chronic respiratory illness or psychiatric illness, or to patients on extracorporeal life support, receiving hyperbaric oxygen therapy, or having elective surgery.

• An upper SpO \textsubscript{2} limit of 96% is therefore reasonable when administering supplemental oxygen to medical patients with acute illness who are not at risk of hypercapnia. However, a higher target may be appropriate for some specific conditions (e.g., pneumothorax, carbon monoxide poisoning, cluster headache, and sickle cell crisis).[116]

• Measure arterial blood gases in those with SpO \textsubscript{2} <94%, those with a risk of hypercapnic ventilatory failure (CO \textsubscript{2} retention), and all patients with high-severity CAP.[1]

Practical tip

Always record the inspired oxygen concentration clearly as this is essential for interpreting blood gas results.

Standard intensive care unit (ICU) supportive care

Arrange for patients with CURB-65 scores of 4 and 5 and an indication for ICU admission to be transferred to ICU and managed by ICU specialists together with respiratory physicians.[1]

• Patients with respiratory failure despite appropriate oxygen therapy require urgent airway management and possible intubation.

Do not routinely give non-invasive ventilation (NIV) or continuous positive airways pressure (CPAP) support in patients with respiratory failure due to CAP.[1]

• If indicated, you should conduct a trial of non-invasive support only in a critical care area where immediate expertise is available to allow a rapid transition to invasive ventilation.[1]

Vasopressors

Start vasopressors if the patient is hypotensive during or after fluid resuscitation to maintain mean arterial pressure level ≥65 mmHg.[117]

Intravenous fluids

Assess all patients for volume depletion and give intravenous fluids if required.[1]

Venous thromboembolism (VTE) prophylaxis

Consider prophylaxis for VTE with a low molecular weight heparin for all patients who are not fully mobile.[1]

Nutritional support

Arrange nutritional support (whether enteral, parenteral, or via nasogastric feeding) for patients with severe CAP who require a prolonged hospital stay.[1]
Community-acquired pneumonia (non COVID-19)

**Treatment**

**Airway clearance**

*Do not treat people with uncomplicated pneumonia with traditional airway clearance techniques routinely.* If needed, offer these patients advice regarding expectoration of sputum.[1]

*Consider airway clearance techniques* if the patient has difficulty expectorating sputum or if they have a pre-existing lung condition.[1]

**Analgesia**

Give simple analgesia (e.g., paracetamol) as appropriate (e.g., for pleuritic pain).[1]

---

**Practical tip**

Encourage patients with uncomplicated CAP (i.e., not complicated by the presence of parapneumonic effusion, empyema, abscess, pneumothorax, necrotising pneumonia, or bronchopleural fistula), whose medical condition allows them to, to sit out of bed. Initially aim for at least 20 minutes in the first 24 hours, and then increase mobility each subsequent day of hospitalisation.

---

**Other treatments**

**Corticosteroids**

*Do not give corticosteroids routinely to patients with CAP of any severity.* [1] [64]

- **Discuss with a senior colleague** patients with comorbidities for which corticosteroids are indicated.[64]

---

**Debate: Corticosteroid treatment in CAP**

*Latest evidence suggests corticosteroids given as an adjunct to antibiotic treatment improve outcomes in adults with CAP but increases the risk of hyperglycaemia. The recommendation from NICE not to give corticosteroids routinely remains valid until further evidence is available.*

- **A Cochrane review** including 17 studies evaluated the safety and efficacy of corticosteroids given as an adjunct to antibiotic treatment for adults and children with pneumonia (CAP, hospital-acquired pneumonia, and ventilator-associated pneumonia). The intervention included oral prednisolone in 3 trials and intravenous dexamethasone, hydrocortisone, or methylprednisolone in 13 trials. The findings suggest that corticosteroids significantly reduced mortality and morbidity in patients with high-severity CAP. They also reduced morbidity, although not mortality, in patients with moderate- or low-severity CAP. Hyperglycaemia was the most common adverse event in adults treated with corticosteroids; however, the authors concluded that overall the benefits outweigh the harms.[118]

- **Two systematic reviews** (published before the Cochrane review) also looked at the safety and efficacy of adjunctive glucocorticoids for patients with CAP.[119] [120] They showed no significant difference in all-cause mortality and reported significant reductions in length of ICU stay,[119] length of hospital stay,[119] [120] and length of time to clinical stability in the corticosteroid groups.[119] [120] One of the reviews reported an increased risk of hyperglycaemia in patients treated with corticosteroids.[120]
• NICE is due to review this area when the results of two ongoing trials have been published. For now, the recommendation not to give corticosteroids routinely remains valid. [64]

**Monitoring**

**Measure observations** initially at least twice daily, and more frequently (e.g., every hour) in **those admitted to a critical care unit (high-dependency unit or ICU)**. [1] **Monitor:**

- Pulse
- Blood pressure
- Respiratory rate
- Temperature
- Oxygen saturation (with a recording of the inspired oxygen saturation at the same time)
- Mental status.

Consider measuring a **baseline C-reactive protein concentration** in patients with CAP on admission to hospital, and repeat the test if clinical progress is uncertain after 48 to 72 hours. [1] [64]

**Review all patients with high-severity CAP** at least every 12 hours until clinical improvement occurs. [1] This should be done by a senior colleague and the medical team. [1]

**Practical tip**

**Pointers to clinical improvement**
The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy: [1]

- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of fever for >24 hours
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphyloccocal, or gram-negative enteric bacilli infection
- No concerns over gastrointestinal absorption.

**Assess severity regularly in all patients** with CAP following hospital admission. [1] An early opportunity for this is the **ward round following admission by a senior colleague and medical team.** [1]

For more information on complications related to CAP see our separate Complications section.

**Failure to improve**

Discuss with a senior colleague any patient who does not improve as expected. [1]

- Consider repeat chest x-ray, C-reactive protein, white cell count, and further specimens for microbiology in patients not progressing satisfactorily after 3 days of treatment. [1]
Community-acquired pneumonia (non COVID-19)

**Treatment**

- Consider **referral to a respiratory physician**.[1]

**Practical tip**

The main reasons why patients **do not improve as expected** include:[1]

- Incorrect diagnosis or complicating condition (e.g., pulmonary embolism, bronchial carcinoma, bronchiectasis)
- Unexpected pathogen or pathogens not covered by antibiotic choice (e.g., ‘atypical’ pathogens, pathogens resistant to commonly used antibiotics such as ampicillin-resistant *Haemophilus influenzae*)
- Antibiotic ineffective or causing allergic reaction (e.g., poor absorption of oral antibiotic, inadequate dose, antibiotic hypersensitivity)
- Impaired local (e.g., bronchiectasis, endobronchial obstruction, aspiration) or systemic (e.g., HIV infection, myeloma) defenses
- Local (e.g., parapneumonic effusion, empyema, lung abscess) or distant (e.g., metastatic infection, sepsicaemia, phlebitis at intravenous cannula site) complications of CAP
- Overwhelming infection
- Improvement expected too soon (e.g., in older patients).

**Discharge from hospital**

**Do not routinely discharge patients if they have had 2* or more** of the following findings present in the **past 24 hours** :[64]

- Temperature >37.5°C (>99.5°F)
- Heart rate >100/minute
- Respiratory rate ≥24/minute
- Systolic blood pressure ≤90 mmHg
- Oxygen saturation <90% on room air
- Inability to maintain oral intake
- Abnormal mental status.

(*The BTS recommends basing your decision to discharge patients with CAP on 1 or more of the findings listed [unless they represent the usual baseline status for that patient] and uses a temperature threshold of 37.8°C [100.04°F]. See **Debate** below.[1])

**Consider delaying discharge** if patients with CAP if **their temperature is higher than 37.5°C (99.5°F).**[64]

**Debate: Cut-off temperature for safe discharge**

*The BTS considers a temperature >37.8°C (>100.04°F), rather than the >37.5°C (>99.5°F) threshold recommended by NICE, as a finding that should prompt you to consider delaying discharge in a patient with CAP.*[1] #

- The >37.5°C (>99.5°F) threshold that is recommended by NICE[64] is based on the conclusions of a prospective cohort study that looked at the value of simple clinical variables for predicting short-term outcomes in patients with pneumonia.[121]
- It found a cut-off of 37.5°C (99.5°F) to be strongly associated with 30-day mortality risk.
Community-acquired pneumonia (non COVID-19)

### Treatment

- For this reason we have based our recommendation on NICE guidance.

At discharge or at follow-up, **offer patients access to information about CAP**, such as a patient information leaflet.[1]

**Arrange a follow-up visit** at around 6 weeks either with the patient’s general practitioner or in a hospital clinic.[1]

#### Follow-up after discharge

**Request a repeat chest x-ray during recovery after about 6 weeks** for patients (regardless of whether they have been admitted to hospital).[1]

- With **persisting symptoms or physical signs**
- Who are at **higher risk of underlying malignancy** (especially smokers and those aged >50 years).

Consider **bronchoscopy during recovery** in patients with **persisting signs, symptoms, and radiological abnormalities at around 6 weeks** after completing treatment.[1]

- Consider a **chest and upper abdomen CT scan** in these patients prior to bronchoscopy.[98]

### Management in the community

Use the **CRB-65 mortality risk tool and your clinical judgement** to determine which patients are suitable for management in the community (see our section above on Risk stratification).

**Consider treatment at home for patients with a CRB-65 score of 0 (low-severity) or a CRB-65 score of 1 or 2 (medium-severity) if they wish to be treated at home and they meet all of the following criteria:** [1] [64]

- They are able to take oral medication safely and reliably
- Their social circumstances make them suitable for treatment at home
- They do not have unstable comorbidities.

**Take a cautious approach, however, when deciding whether it is safe to treat any patient with moderate-severity CAP (CRB-65 score of 1 or 2) in the community.** These patients are at increased risk of death, particularly those with a CRB-65 score of 2. **You should refer the majority for management in hospital.** [1] [64]

- If you decide to treat the patient in the community, follow the same treatment recommendations given below for patients with suspected CAP presenting in the community (low-severity).

**Refer any patients presenting in the community with a CRB-65 score of 3 or more for immediate hospital admission (usually by blue-light ambulance in the UK).** [1] [64]

### Antibiotics

The following recommendations are **based on guidelines from the British Thoracic Society (BTS) and the National Institute for Health and Care Excellence (NICE)**.[1] [105]
Giving antibiotics prior to hospital transfer

High-severity CAP (CRB-65 score 3 or 4)

Give empirical antibiotics prior to hospital transfer (usually by blue-light ambulance in the UK) to patients with suspected CAP considered to be life-threatening. Follow your local protocol.

- The first-line choice is intravenous benzylpenicillin or oral amoxicillin. Clarithromycin is an alternative for people who are allergic to penicillin.

- More recent (2019) guidelines from NICE on antimicrobial prescribing in adults recommend amoxicillin/clavulanate plus clarithromycin (or erythromycin) first-line in people with high-severity CAP. These recommendations are based mainly on expert opinion as the evidence is limited.

Consider giving antibiotics prior to hospital transfer to patients with suspected high-severity CAP where there are likely to be delays of over 6 hours to hospital admission and treatment.

- Pre-admission antibiotics can negatively influence the results of subsequent microbiological investigations, but this is not seen as a reason for withholding antibiotics if a general practitioner considers it indicated.

Moderate-severity CAP (CRB-65 score 1 or 2) and low-severity CAP (CRB-65 score 0)

Do not give antibiotics to patients with moderate-severity CAP or low-severity CAP prior to hospital referral.

Empirical antibiotics for home treatment

Give empirical oral antibiotics to patients treated in the community.

- The first-line option is amoxicillin. [1][105]

- Alternative options for patients who are allergic to penicillin are a macrolide (e.g., clarithromycin) or a tetracycline (e.g., doxycycline).

- NICE recommends clarithromycin (or erythromycin) or doxycycline as alternatives to amoxicillin for patients allergic to penicillin and for patients in whom amoxicillin is unsuitable (e.g., if atypical pneumonia is suspected). NICE recommendations are based mainly on expert opinion as the evidence is limited.

Consider adding or switching to a macrolide if the patient does not respond to amoxicillin.

- Consider whether a correct diagnosis of CAP has been made if a patient does not respond to initial empirical antibiotics.

- Consider clinical and radiographic review to look for secondary diagnoses or complications of CAP such as pleural effusion/empyema, lung abscess, or worsening pneumonic shadowing.
• Consider changing initial empirical antibiotics but first consider compliance with and adequate absorption of an oral regimen.[1]

Give antibiotic therapy for 5 days. NICE recommends :[105]

• Prescribing the shortest course that is likely to be effective to reduce the risk of antimicrobial resistance and minimise the risk of adverse effects
• Stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable. This should be based on your clinical judgement and the following criteria:[105] [112]

• Fever in past 48 hours, or more than one sign of clinical instability:
  • Systolic blood pressure <90 mmHg
  • Heart rate >100/minute
  • Respiratory rate >24/minute
  • Arterial oxygen saturation <90% or PaO$_2$ <60 mmHg in room air.

General management
Advise patients to rest, to drink plenty of fluids, and not to smoke.[1]

Failure to respond
Advise patients (and their carers) to seek medical advice if:[105]

• Symptoms worsen rapidly or significantly
• Symptoms do not start to improve within 3 days
• The person becomes systemically very unwell.

Around 10% of patients managed in the community do not respond to antibiotic therapy and require hospitalisation.[122]

Admit urgently to hospital any patient on antibiotic treatment with features of moderate- or high-severity infection .[1]

Follow-up
Order a chest x-ray during recovery after about 6 weeks for patients (regardless of whether patients have been admitted to hospital).[1]

• With persisting symptoms or physical signs
• Who are at higher risk of underlying malignancy (especially people who smoke and those aged >50 years).

Consider bronchoscopy during recovery in patients with persisting signs, symptoms, and radiological abnormalities at around 6 weeks after completing treatment .[1]

• Consider a chest and upper abdomen CT scan in these patients prior to bronchoscopy.[98]
Prognosis

For patients admitted to hospital, mortality rate ranges from 5% to 15%, but increases to 20% to 50% in patients requiring admission to the intensive care unit (ICU).[6] [106]

- Risk factors associated with increased 30-day mortality include bacteraemia, admission to the ICU, comorbidities (especially neurological disease), and infection with a potentially multidrug-resistant pathogen (e.g., *Staphylococcus aureus*, *Pseudomonas aeruginosa*, Enterobacteriaceae).[36] [123] [124] [125]

Patients treated in the community generally have a good prognosis.[1]

Treatment details overview

Please note that formulations/routes and doses may differ between drug names and brands, drug formularies, or locations. Treatment recommendations are specific to patient groups: see disclaimer

<table>
<thead>
<tr>
<th>Initial</th>
<th>( summary )</th>
</tr>
</thead>
<tbody>
<tr>
<td>suspected CAP: presenting in hospital with life-threatening illness</td>
<td>1st empirical intravenous antibiotic therapy plus supportive care</td>
</tr>
<tr>
<td>suspected CAP: presenting in hospital without life-threatening illness</td>
<td>1st supportive care while confirming diagnosis</td>
</tr>
<tr>
<td>suspected CAP: presenting in the community</td>
<td>1st urgent hospital admission</td>
</tr>
<tr>
<td>high-severity (CRB-65 = 3 or 4)</td>
<td>consider empirical antibiotic prior to hospital transfer</td>
</tr>
<tr>
<td>moderate-severity (CRB-65 = 1 or 2)#</td>
<td>1st hospital referral</td>
</tr>
<tr>
<td>low-severity (CRB-65 = 0)</td>
<td>1st empirical oral antibiotic therapy plus supportive care consider hospital admission</td>
</tr>
</tbody>
</table>
## Acute

<table>
<thead>
<tr>
<th>confirmed CAP on chest x-ray: presenting in hospital</th>
</tr>
</thead>
<tbody>
<tr>
<td>high-severity (CURB-65 = 3-5)</td>
</tr>
<tr>
<td>1st empirical intravenous antibiotic therapy</td>
</tr>
<tr>
<td>consider fluoroquinolone</td>
</tr>
<tr>
<td>plus supportive care</td>
</tr>
<tr>
<td>consider switch to pathogen-targeted antibiotic therapy</td>
</tr>
<tr>
<td>moderate-severity (CURB-65 = 2)</td>
</tr>
<tr>
<td>1st empirical oral or intravenous antibiotic therapy</td>
</tr>
<tr>
<td>plus supportive care</td>
</tr>
<tr>
<td>consider switch to pathogen-targeted antibiotic therapy</td>
</tr>
<tr>
<td>low-severity (CURB-65 = 0-1)</td>
</tr>
<tr>
<td>1st empirical oral or intravenous antibiotic therapy</td>
</tr>
<tr>
<td>plus supportive care</td>
</tr>
<tr>
<td>consider switch to pathogen-targeted antibiotic therapy</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>confirmed CAP on chest x-ray: presenting in the community</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st continue empirical antibiotics or switch to pathogen-targeted antibiotic therapy</td>
</tr>
</tbody>
</table>
Treatment options

Please note that formulations/routes and doses may differ between drug names and brands, drug formularies, or locations. Treatment recommendations are specific to patient groups: see disclaimer.
### Initial

suspected CAP: presenting in hospital with life-threatening illness

1st  empirical intravenous antibiotic therapy

» Give empirical antibiotics to patients presenting in hospital with life-threatening disease based on a presumptive clinical diagnosis of CAP.

  • Follow your local protocol (e.g., Sepsis Six or Surviving Sepsis Campaign 1-hour care bundle) for investigating and treating all patients with suspected sepsis, or those at risk, within 1 hour.\[66\] [67] [68]

  » Immediately order a chest x-ray to confirm the diagnosis. [1]

  • Once a diagnosis of CAP is confirmed, manage these patients as per the protocols below for patients with confirmed CAP on chest x-ray: presenting in hospital.

plus  supportive care

Treatment recommended for ALL patients in selected patient group

**Primary options**

» **paracetamol**: 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/day; 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day

» **Provide supportive care**, which may include the following measures.[1]

» **Oxygen**

  » Prescribe oxygen if oxygen saturation <94% and maintain at target range.[1] For patients at risk of CO\textsubscript{2} retention prescribe oxygen if oxygen saturation <88%.

  • **Target oxygen saturation range of**:  
    - 94% to 96% in acutely ill patients who are **not at risk of hypercapnia** [72]
    - 88% to 92% in patients **at risk of hypercapnia** [71]
    - **Measure arterial blood gases** in those with SpO\textsubscript{2} <94%, those with a risk of hypercapnic ventilatory failure (CO2
<table>
<thead>
<tr>
<th><strong>Initial</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial retention), and all patients with high-severity CAP. [1]</td>
</tr>
</tbody>
</table>

**Practical tip**

Always record the inspired oxygen concentration clearly as this is essential for interpreting blood gas results.

»

» **Fluid resuscitation**

- Assess all patients for volume depletion and give intravenous fluids if required. [1]

»

» **Standard intensive care unit (ICU) supportive care**

- Arrange for patients with an indication for ICU admission to be transferred to ICU and managed by ICU specialists together with respiratory physicians. [1]

- Patients with respiratory failure despite appropriate oxygen therapy require urgent airway management and possible intubation.

- Do not routinely give non-invasive ventilation (NIV) or continuous positive airways pressure (CPAP) support in patients with respiratory failure due to CAP. [1]

  - If indicated, you should conduct a trial of non-invasive support only in a critical care area where immediate expertise is available to allow a rapid transition to invasive ventilation. [1]

» **Vasopressors**

- Start vasopressors if the patient is hypotensive during or after fluid resuscitation to maintain mean arterial pressure level greater than or equal to 65 mmHg. [117] See our Sepsis topic for more information.

» **Analgesia**

- Give simple analgesia as appropriate (e.g., for pleuritic pain). [1]
### Initial

**suspected CAP: presenting in hospital without life-threatening illness**

1st support care while confirming diagnosis

**Primary options**

- **paracetamol:** 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/day; 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day

- Confirm diagnosis by chest x-ray before starting antibiotic therapy.

- In patients presenting in hospital without life-threatening illness, confirm the diagnosis by chest x-ray before starting antibiotics. Once diagnosis is confirmed, patients are managed as per the protocols below for patients with confirmed CAP on chest x-ray: presenting in hospital.

- In the meantime, provide supportive care as necessary, which may include the following measures. Oxygen

- Prescribe oxygen if oxygen saturation <94% and maintain at target range. For patients at risk of hypercapnia prescribe oxygen if oxygen saturation <88%.

- Target oxygen saturation range of:
  - 94% to 96% in acutely ill patients who are not at risk of hypercapnia [72]
  - 88% to 92% in patients at risk of hypercapnia [71]

- Measure arterial blood gases in those with SpO₂ <94%, those with a risk of hypercapnic ventilatory failure (CO₂ retention), and all patients with high-severity CAP.[1]

**Practical tip**

Always record the inspired oxygen concentration clearly as this is essential for interpreting blood gas results.
Initial

» Fluid resuscitation

- Assess all patients for volume depletion and give intravenous fluids if required.[1]

» Standard intensive care unit (ICU) supportive care

- Arrange for patients with an indication for ICU admission to be transferred to ICU and managed by ICU specialists together with respiratory physicians.[1]

- Patients with respiratory failure despite appropriate oxygen therapy require urgent airway management and possible intubation.

- Do not routinely give non-invasive ventilation (NIV) or continuous positive airways pressure (CPAP) support in patients with respiratory failure due to CAP.[1]

- If indicated, you should conduct a trial of non-invasive support only in a critical care area where immediate expertise is available to allow a rapid transition to invasive ventilation.[1]

» Vasopressors

- Start vasopressors if the patient is hypotensive during or after fluid resuscitation to maintain mean arterial pressure level greater than or equal to 65 mmHg.[17] See our Sepsis topic for more information.

» Analgesia

- Give simple analgesia as appropriate (e.g., for pleuritic pain).[1]

suspected CAP: presenting in the community

- high-severity (CRB-65 = 3 or 4)

1st urgent hospital admission

- Refer patients presenting in the community with high-severity CAP (CRB-65 score of 3 or 4) for immediate hospital admission (usually by blue-light ambulance in the UK). [1] [64]
### Initial

<table>
<thead>
<tr>
<th>moderate-severity (CRB-65 = 1 or 2)#</th>
<th>1st hospital referral</th>
</tr>
</thead>
<tbody>
<tr>
<td>• In hospital, once the diagnosis of CAP is confirmed by chest x-ray and the disease severity has been assessed, patients are managed as per the protocols below for patients with confirmed CAP on chest x-ray: presenting in hospital.</td>
<td>• Refer patients presenting in the community with moderate-severity CAP (CRB-65 score of 1 or 2) to hospital for assessment and management.# These patients are at increased risk of death, particularly those with a score of 2.[1] [64]</td>
</tr>
<tr>
<td>consider empirical antibiotic prior to hospital transfer</td>
<td>• Pre-admission antibiotics can negatively influence the results of subsequent microbiological investigations, but this is not seen as a reason for withholding antibiotics if a general practitioner considers it indicated.[1]</td>
</tr>
<tr>
<td>Treatment recommended for SOME patients in selected patient group</td>
<td>» Consult your local protocol for guidance on selection of antibiotic regimen.</td>
</tr>
<tr>
<td>» Give empirical antibiotics prior to hospital transfer (usually by blue-light ambulance in the UK) to any patients with suspected high-severity CAP considered to be life-threatening, according to your local protocol.[1]</td>
<td></td>
</tr>
<tr>
<td>• British Thoracic Society guidelines recommend intravenous benzylpenicillin or oral amoxicillin. Oral clarithromycin is an alternative for people who are allergic to penicillin.[1]</td>
<td></td>
</tr>
<tr>
<td>» Consider giving empirical antibiotics prior to hospital transfer to patients with suspected high-severity CAP where there are likely to be delays of over 6 hours to hospital admission and treatment.[1]</td>
<td></td>
</tr>
</tbody>
</table>
**Initial**

» **Consider managing patients in the community** if they prefer to be treated at home and they meet the following criteria:[1] [64]

- They are able to take oral medication safely and reliably
- Their social circumstances make them suitable for treatment at home
- They do not have unstable comorbidities.

» **Take a cautious approach, however, when deciding whether it is safe to treat any patient with moderate-severity CAP in the community.** You should refer the majority for management in hospital.[1] [64]

- If you decide to treat the patient in the community, follow the same treatment recommendations given below for patients with suspected CAP: presenting in the community (low-severity).

**low-severity (CRB-65 = 0)**

1st empirical oral antibiotic therapy

» **Give empirical oral antibiotics and manage people with low-severity CAP (CRB-65 score of 0) in the community.** [1] [64]

- The first-line option is amoxicillin.
  [1] Alternative options for patients who are **allergic to penicillin** are a macrolide (e.g., clarithromycin) or a tetracycline (e.g., doxycycline).[1]
- If the patient does not respond to amoxicillin monotherapy, consider adding, or switching to, a macrolide (e.g., clarithromycin).[1]

» Advise patients (and their carers) to **seek medical advice** if their symptoms worsen rapidly or significantly, their symptoms do not start to improve within 3 days, or they become systemically very unwell.[105]

- Admit urgently to hospital any patient on antibiotic treatment with features of **moderate- or high-severity infection**.[1] Around 10% of patients managed in the community do not respond to antibiotic therapy and require hospitalisation.[122]

» **Give antibiotic treatment for 5 days.**[105] The National Institute for Health and Care Excellence recommends **stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is**
not clinically stable. This should be based on your clinical judgement and the following criteria:[1] [112]

- Fever in past 48 hours, or more than one sign of clinical instability:
  - Systolic blood pressure <90 mmHg
  - Heart rate >100/minute
  - Respiratory rate >24/minute
  - Arterial oxygen saturation <90% or PaO₂ <60 mmHg in room air.

» Consult local protocols for guidance on selection of antibiotic regimen.

plus supportive care

Treatment recommended for ALL patients in selected patient group

Primary options

» paracetamol: 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/day; 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day

» Advise patients to rest, to drink plenty of fluids, and not to smoke.[1]

» Give simple analgesia as appropriate (e.g., for pleuritic pain).[1]

consider hospital admission

Treatment recommended for SOME patients in selected patient group

» Consider referring patients to hospital if:[1] [64]

- They are not able to take oral medication safely and reliably
- Their social circumstances do not make them suitable for treatment at home
- They have unstable comorbidities
- They prefer to be treated in hospital.

» In hospital, once the diagnosis of CAP is confirmed by chest x-ray and the disease severity has been assessed, patients are managed as per the protocols below for patients with confirmed CAP on chest x-ray: presenting in hospital.
**Acute**

**confirmed CAP on chest x-ray: presenting in hospital**

- high-severity (CURB-65 = 3-5)  
  1st empirical intravenous antibiotic therapy

  » Always manage patients with high-severity CAP in hospital. [1]

  » Give empirical broad-spectrum intravenous antibiotics immediately after diagnosis. This should be within 4 hours of presentation to hospital. [1] [105]

  » Prescribe an appropriate antibiotic regimen according to your local protocol to help reduce the development of antibiotic resistance and *Clostridium difficile* infection. Consult with a microbiologist. The British Thoracic Society (BTS) recommends:[1]

  - A broad-spectrum beta-lactamase-resistant penicillin (e.g., amoxicillin/clavulanate) plus a macrolide (e.g., clarithromycin).[1]
  - For patients who are allergic to penicillin, give a second-generation cephalosporin (e.g., cefuroxime) or a third-generation cephalosporin (e.g., cefotaxime or ceftriaxone) plus a macrolide (e.g., clarithromycin).[1]
  - A small number of patients are allergic to both penicillins and cephalosporins; consult an infectious disease consultant for selection of appropriate antibiotics in these patients.

  » Review route of administration initially on the ward round following admission and then daily thereafter. [1] De-escalate treatment as soon as appropriate, including switching from intravenous to oral therapy. [1] When making this decision consider response to treatment (see practical tip), change in disease severity, and contraindications to oral administration such as:[1]

  - Patient is unable to swallow (e.g., impaired swallowing reflex, impaired consciousness)
  - Gastrointestinal malabsorption for functional or anatomical reasons.

**Practical tip**

**Pointers to clinical improvement**

---

This PDF of the BMJ Best Practice topic is based on the web version that was last updated: Jun 04, 2020. BMJ Best Practice topics are regularly updated and the most recent version of the topics can be found on bestpractice.bmj.com. Use of this content is subject to our disclaimer. © BMJ Publishing Group Ltd 2020. All rights reserved.
Acute

The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]

- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of fever for >24 hours
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
- No concerns over gastrointestinal absorption.

» Give antibiotic therapy for 5 days. [105] The National Institute for Health and Care Excellence recommends stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable. [105] This should be based on your clinical judgement and the following criteria:[105] [112]

- Fever in past 48 hours, or more than one sign of clinical instability:
  - Systolic blood pressure <90 mmHg
  - Heart rate >100/minute
  - Respiratory rate >24/minute
  - Arterial oxygen saturation <90% or PaO₂ <60 mmHg in room air.

» In some people, longer courses might be needed due to individual circumstances. In the UK, some hospitals require consultation with the microbiology team if considering extending the duration of antibiotic treatment beyond 5 days in high-severity CAP. Follow your local protocol.

» Consult your local protocol for guidance on selection of antibiotic regimen.

consider fluoroquinolone

Treatment recommended for SOME patients in selected patient group
### Acute

- **Consult with a microbiologist and senior clinician before giving a fluoroquinolone.**
  - Consider safety issues associated with fluoroquinolone use. Fluoroquinolones are known to cause tendonitis, tendon rupture, arthralgia, neuropathies, and other musculoskeletal or nervous system effects.\(^{[113]}\) \(^{[114]}\)
  - The British Thoracic Society guideline recommends adding a fluoroquinolone to the existing empirical regimen (i.e., triple therapy) if the patient does not respond, or if legionella pneumonia is strongly suspected.\(^{[1]}\) However, in practice there are concerns about the risk of using a macrolide and a fluoroquinolone together as they can both prolong the QT interval. Therefore, some clinicians may replace the macrolide in the original empirical regimen with a fluoroquinolone instead (i.e., dual therapy).

**plus supportive care**

Treatment recommended for ALL patients in selected patient group

**Primary options**

- **paracetamol**: 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/day; 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day

- **Oxygen**
  - Prescribe oxygen if oxygen saturation <94% and maintain at target range.\(^{[1]}\) For patients at risk of hypercapnia prescribe oxygen if oxygen saturation <88%.
  - Target oxygen saturation range of:
    - **94% to 96%** in acutely ill patients who are not at risk of hypercapnia \(^{[72]}\)
    - **88% to 92%** in patients at risk of hypercapnia \(^{[71]}\)
  - Measure arterial blood gases in those with \(\text{SpO}_2\) <94%, those with a risk of hypercapnic ventilatory failure (\(\text{CO}_2\)
### Acute

- **Community-acquired pneumonia (non COVID-19)**

  Treatment

  Acute retention, and all patients with high-severity CAP.[1]

  - **Fluid resuscitation**

    - Assess all patients for volume depletion and give intravenous fluids if required.[1]

  - **Standard intensive care unit (ICU) supportive care**

    - Arrange for patients with CURB-65 scores of 4 and 5 and an indication for ICU admission to be transferred to ICU and managed by ICU specialists together with respiratory physicians.[1]

    - Patients with respiratory failure despite appropriate oxygen therapy require urgent airway management and possible intubation.

    - Do not routinely give non-invasive ventilation (NIV) or continuous positive airways pressure (CPAP) support in patients with respiratory failure due to CAP.[1]

      - If indicated, you should conduct a trial of non-invasive support only in a critical care area where immediate expertise is available to allow a rapid transition to invasive ventilation.[1]

  - **Vasopressors**

    - Start vasopressors if the patient is hypotensive during or after fluid resuscitation to maintain mean arterial pressure level greater than or equal to 65 mmHg.[117]

  - **Venous thromboembolism (VTE) prophylaxis**

    - Consider prophylaxis for VTE with a low molecular weight heparin for all patients who are not fully mobile.[1] In practice in the UK, prescription of heparin will be prompted if appropriate once you have recorded your VTE risk assessment in the patient’s electronic record.

  - **Nutritional support**
### Acute

- Arrange nutritional support (whether enteral, parenteral, or via nasogastric feeding) for patients with severe CAP who require a prolonged hospital stay.\(^1\)
  - **Airway clearance**
    - Do not treat people with uncomplicated pneumonia with traditional airway clearance techniques routinely. If needed, offer these patients advice regarding expectoration of sputum.\(^1\)
    - Consider airway clearance techniques if the patient has difficulty expectorating sputum or if they have a pre-existing lung condition.\(^1\)
  - **Analgesia**
    - Give simple analgesia as appropriate (e.g., for *pleuritic pain*).\(^1\)

### Consider switch to pathogen-targeted antibiotic therapy

Treatment recommended for SOME patients in selected patient group

- Consult with a microbiologist about appropriate pathogen-targeted antibiotic therapy.\(^1\)
  - Switch from empirical antibiotics to pathogen-targeted antibiotics as soon as specific pathogens are identified (unless there are legitimate concerns about dual-pathogen infection).\(^1\)
  - Only about one third to one quarter of patients with CAP admitted to hospital will have their pneumonia defined microbiologically.\(^1\) Among these patients:
    - Around 14% have an atypical pathogen, of which:\(^{21}\)
      - 7% have *Mycoplasma pneumoniae*
      - 4% have *Chlamydophila pneumoniae*
      - 3% have *Legionella pneumophila*.
    - Those with infections due to *Mycoplasma*, *Chlamydophila*, and *Coxiella burnetii* will be diagnosed late in the illness on the basis of seroconversion, reducing the opportunity for early targeted therapy.\(^1\)
    - Consider switching the choice of agent once the results of sensitivity testing are available.
Community-acquired pneumonia (non COVID-19)

Treatment

or following consultation with a microbiologist, intensivist, or respiratory physician.[1]

» BTS recommendations for pathogen-targeted antibiotics [1]

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Mycoplasma pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia pneumoniae</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Legionella species</em></td>
<td>Fluoroquinolone (orally or intravenously)</td>
<td>Clarithromycin (orally or intravenously) or azithromycin (in countries where it is used for managing pneumonia)</td>
</tr>
<tr>
<td><em>Streptococcus pneumoniae</em></td>
<td>Amoxicillin (orally) or benzylpenicillin (intravenously)</td>
<td>Clarithromycin (orally) or cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia psittaci</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Coxiella burnetii</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Haemophilus influenzae</em></td>
<td>Non-beta-lactamase-producing : amoxicillin (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td></td>
<td>Beta-lactamase-producing : amoxicillin/clavulanate</td>
<td></td>
</tr>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
<td>Alternative antibiotic</td>
</tr>
<tr>
<td>----------------------------------</td>
<td>----------------------</td>
<td>------------------------</td>
</tr>
<tr>
<td></td>
<td>(orally or intravenously)</td>
<td>Fluoroquinolone (intravenously) or imipenem/cilastatin (intravenously) or meropenem (intravenously)</td>
</tr>
<tr>
<td>Gram-negative enteric bacilli</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
<td>Ciprofloxacin (intravenously) or piperacillin/tazobactam (intravenously) plus Gentamicin or tobramycin (dose monitoring required)</td>
</tr>
<tr>
<td>\textit{Pseudomonas aeruginosa}</td>
<td>Ceftazidime (intravenously) plus Gentamicin or tobramycin (dose monitoring required)</td>
<td>Vancomycin (intravenously; dose monitoring required) or linezolid (intravenously) or teicoplanin (intravenously) with or without</td>
</tr>
<tr>
<td>\textit{Staphylococcus aureus} : non-MRSA</td>
<td>Flucloxacillin (intravenously) with or without Rifampicin (orally or intravenously)</td>
<td>\textit{Staphylococcus aureus} : MRSA</td>
</tr>
</tbody>
</table>
Community-acquired pneumonia (non COVID-19)

Treatment

<table>
<thead>
<tr>
<th>Acute</th>
<th>1st empirical oral or intravenous antibiotic therapy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
</tr>
<tr>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
</tbody>
</table>

» Consider patients with moderate-severity CAP for short-stay inpatient treatment or hospital-supervised outpatient treatment. [1]

» Give antibiotics as soon as possible after diagnosis. This should be within 4 hours of presentation to hospital. [1] [105]

» Give broad-spectrum empirical oral antibiotics.

- Most patients with moderate-severity CAP can be treated with dual oral antibiotic therapy. [1] British Thoracic Society guidelines recommend amoxicillin plus a macrolide (e.g., clarithromycin). [1]

- For patients who are allergic to penicillin or macrolides, consider oral doxycycline. [1] Alternative choices include oral levofloxacin or moxifloxacin (after considering safety issues associated with fluoroquinolone use). [1]

- If oral antibiotics are contraindicated (e.g., patient is unable to swallow or has gastrointestinal malabsorption for functional or anatomical reasons) give intravenous amoxicillin or benzylpenicillin plus clarithromycin. [1]

- For patients who are allergic to penicillin in whom oral antibiotics are contraindicated, give a second-generation cephalosporin (e.g., cefuroxime) or a third-generation cephalosporin (e.g., cefotaxime or ceftriaxone) plus clarithromycin, or intravenous levofloxacin monotherapy. [1]

- If the patient does not respond to a combination of amoxicillin and clarithromycin, consider changing treatment to doxycycline or a fluoroquinolone with effective pneumococcal cover (e.g., levofloxacin, moxifloxacin). [1]
### Acute

**More info: EMA and MHRA restrictions on the use of fluoroquinolone antibiotics**

In November 2018, the European Medicines Agency (EMA) completed a review of serious, disabling, and potentially irreversible adverse effects associated with systemic and inhaled fluoroquinolone antibiotics. These adverse effects include tendonitis, tendon rupture, arthralgia, neuropathies, and other musculoskeletal or nervous system effects.

- As a consequence of this review, the EMA now recommends that fluoroquinolone antibiotics be restricted for use in serious, life-threatening bacterial infections only. Furthermore, they recommend that fluoroquinolones should not be used for mild to moderate infections unless other appropriate antibiotics for the specific infection cannot be used, and should not be used in non-severe, non-bacterial, or self-limiting infections. Patients who are older, have renal impairment, or have had a solid organ transplant, and those being treated with a corticosteroid are at a higher risk of tendon damage. Co-administration of a fluoroquinolone and a corticosteroid should be avoided. The UK-based Medicines and Healthcare products Regulatory Agency (MHRA) supports these recommendations.

- For this reason, fluoroquinolones (e.g., levofloxacin, moxifloxacin) should only be considered in moderate-severity CAP when it is considered inappropriate to use other antibiotics that are commonly recommended for the treatment of CAP. **Consult with a microbiologist about whether a fluoroquinolone is an appropriate option for your patient.**

- Consider monotherapy with a macrolide for patients who have been treated in the community and who have not responded to an adequate course of amoxicillin prior to hospital admission.

- Deciding whether the course of amoxicillin was adequate is tricky and involves...
## Acute

Clinical judgement. **Consult a senior clinician before prescribing monotherapy within the first 24 hours of admission.**[1]**  

> Review the need for intravenous antibiotics initially on the ward round following admission and then every day thereafter.[1]**  

- Switch to oral antibiotics as soon as clinical improvement occurs (see practical tip), and as long as there are no contraindications to oral administration (e.g., patient is unable to swallow or has gastrointestinal malabsorption for functional or anatomical reasons).[1]

### Practical tip

**Pointers to clinical improvement**

The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]

- Pulse rate <100 beats/minute
- Resolution of tachypnoea
- Clinically hydrated and taking oral fluids
- Resolution of fever for >24 hours
- Resolution of hypotension
- Absence of hypoxia
- Improving white cell count
- Non-bacteraemic infection
- No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
- No concerns over gastrointestinal absorption.

> **Give antibiotic therapy for 5 days.**[105] The National Institute for Health and Care Excellence recommends **stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable.**[105] This should be based on your clinical judgement and the following criteria:[1][112]

- Fever in past 48 hours, or more than one sign of clinical instability:
  - Systolic blood pressure <90 mmHg
  - Heart rate >100/minute
  - Respiratory rate >24/minute
Community-acquired pneumonia (non COVID-19)

### Treatment

<table>
<thead>
<tr>
<th>Acute</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Arterial oxygen saturation &lt;90% or PaO&lt;sub&gt;2&lt;/sub&gt; &lt; 60 mmHg in room air.</td>
</tr>
</tbody>
</table>

» **In some people, longer courses might be needed due to individual circumstances.** In the UK, some hospitals require consultation with the microbiology team if considering extending the duration of antibiotic treatment beyond 5 days in moderate-severity CAP. **Follow your local protocol.**

» Consult your local protocol for guidance on selection of antibiotic regimen.

**plus** **supportive care**

Treatment recommended for ALL patients in selected patient group

**Primary options**

» **paracetamol**: 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/day; 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day

» Provide supportive care, which may include the following measures.[1]

» **Oxygen**

» Prescribe oxygen if oxygen saturation <94% and maintain at target range.[1] For patients at risk of hypercapnia prescribe oxygen if oxygen saturation <88%.

• **Target oxygen saturation range of**:  
  • 94% to 96% in acutely ill patients who are **not at risk of hypercapnia**.[72]  
  • 88% to 92% in patients **at risk of hypercapnia**.[71]

• **Measure arterial blood gases** in those with SpO<sub>2</sub> < 94%, those with a risk of hypercapnic ventilatory failure (CO<sub>2</sub> retention), and all patients with high-severity CAP.[1]

»

» **Fluid resuscitation**

• Assess all patients for volume depletion and give intravenous fluids if required.[1]

»

» **Venous thromboembolism (VTE) prophylaxis**
Community-acquired pneumonia (non COVID-19)

**Acute**

- Consider prophylaxis for VTE with a low molecular weight heparin for all patients who are not fully mobile.[1]

  » **Airway clearance**

  - Do not treat people with uncomplicated pneumonia with traditional airway clearance techniques routinely. If needed, offer these patients advice regarding expectoration of sputum.[1]
  - Consider airway clearance techniques if the patient has difficulty expectorating sputum or if they have a pre-existing lung condition.[1]

  » **Analgesia**

  - Give simple analgesia as appropriate (e.g., for pleuritic pain).[1]

**consider switch to pathogen-targeted antibiotic therapy**

Treatment recommended for SOME patients in selected patient group

- **Consult with a microbiologist about appropriate pathogen-targeted antibiotic therapy.**[1]

  - Switch from empirical antibiotics to pathogen-targeted antibiotics as soon as specific pathogens are identified (unless there are legitimate concerns about dual-pathogen infection).[1]

  » Only about one third to one quarter of patients with CAP admitted to hospital will have their pneumonia defined microbiologically.[1] Among these patients:

    - Around 14% have an atypical pathogen, of which[21]
      - 7% have *Mycoplasma pneumoniae*
      - 4% have *Chlamydophila pneumoniae*
      - 3% have *Legionella pneumophila*
    - Those with infections due to *Mycoplasma*, *Chlamydophila*, and *Coxiella burnetii* will be diagnosed late in the illness on the basis of seroconversion, reducing the opportunity for early targeted therapy.[1]

  » Consider switching the choice of agent once the results of sensitivity testing are available or following consultation with a microbiologist, intensivist, or respiratory physician.[1]
### Acute

> **BTS recommendations for pathogen-targeted antibiotics** [1]

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Mycoplasma pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Legionella species</em></td>
<td>Fluoroquinolone (orally or intravenously)</td>
<td>Clarithromycin (orally or intravenously) or azithromycin (in countries where it is used for managing pneumonia)</td>
</tr>
<tr>
<td><em>Streptococcus pneumoniae</em></td>
<td>Amoxicillin (orally) or benzylpenicillin (intravenously)</td>
<td>Clarithromycin (orally) or cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia psittaci</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Coxiella burnetii</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Haemophilus influenzae</em></td>
<td>Non-beta-lactamase-producing: amoxicillin (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td></td>
<td>Beta-lactamase-producing: amoxicillin/clavulanate (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
<td>Alternative antibiotic</td>
</tr>
<tr>
<td>----------------------------------</td>
<td>----------------------------------------------</td>
<td>----------------------------------------</td>
</tr>
<tr>
<td><strong>Gram-negative enteric bacilli</strong></td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
<td>Fluoroquinolone (intravenously) or imipenem/cilastatin or meropenem (intravenously)</td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em></td>
<td>Ceftazidime (intravenously) plus Gentamicin or tobramycin (dose monitoring required)</td>
<td>Ciprofloxacin (intravenously) or piperacillin/tazobactam (intravenously) plus Gentamicin or tobramycin (dose monitoring required)</td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em>: non-MRSA</td>
<td>Flucloxacillin (intravenously) with or without Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em>: MRSA</td>
<td>Vancomycin (intravenously; dose monitoring required) or linezolid (intravenously) or teicoplanin (intravenously) with or without</td>
<td></td>
</tr>
</tbody>
</table>
# Community-acquired pneumonia (non COVID-19)

## Treatment

### Acute

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### low-severity (CURB-65 = 0-1)

1st empirical oral or intravenous antibiotic therapy

» Most patients with low-severity CAP can be discharged for treatment at home. However, consider admitting patients if:[1] [64]

- They are not able to take oral medication safely and reliably
- Their social circumstances do not make them suitable for treatment at home
- They have unstable comorbidities
- They prefer to be treated in hospital.

» Give antibiotics as soon as possible. This should be within 4 hours of presentation to hospital. [1] [64]

» Most patients with low-severity CAP managed in hospital can be treated with oral antibiotics.[1]

- The preferred choice is amoxicillin. [1] Consider a macrolide (e.g., clarithromycin) or a tetracycline (e.g., doxycycline) for patients who are allergic to penicillin.[1]

- If the patient does not respond to amoxicillin monotherapy, consider adding, or switching to, a macrolide (e.g., clarithromycin).[1]

- If the oral route is contraindicated (e.g., impaired swallowing reflex, impaired consciousness, gastrointestinal malabsorption) consider intravenous amoxicillin, benzylpenicillin, or clarithromycin.[1]

- Review the need for intravenous antibiotics initially during the ward round following admission and then every day after. [1]

  - Switch to oral antibiotics as soon as clinical improvement occurs (see practical tip), and as long as there are
Community-acquired pneumonia (non COVID-19)

Treatment

Acute

no contraindications to oral administration.

Practical tip

Pointers to clinical improvement
The following clinical features should prompt you to consider switching from intravenous to oral antibiotic therapy:[1]

• Pulse rate <100 beats/minute
• Resolution of tachypnoea
• Clinically hydrated and taking oral fluids
• Resolution of fever for >24 hours
• Resolution of hypotension
• Absence of hypoxia
• Improving white cell count
• Non-bacteraemic infection
• No microbiological evidence of legionella, staphylococcal, or gram-negative enteric bacilli infection
• No concerns over gastrointestinal absorption.

» Give antibiotic therapy for 5 days.[105] The National Institute for Health and Care Excellence recommends stopping treatment after 5 days unless microbiological results suggest a longer course or the patient is not clinically stable.[105] This should be based on your clinical judgement and the following criteria:[105][112]

• Fever in past 48 hours, or more than one sign of clinical instability:
  • Systolic blood pressure <90 mmHg
  • Heart rate >100/minute
  • Respiratory rate >24/minute
  • Arterial oxygen saturation <90% or PaO_2 <60 mmHg in room air.

» Consult local protocols for guidance on selection of antibiotic regimen.

plus supportive care
Treatment recommended for ALL patients in selected patient group

Primary options

» paracetamol: 500-1000 mg orally every 4-6 hours when required, maximum 4000 mg/
**Acute**

| Treatment | 15 mg/kg (maximum 1000 mg/dose) intravenously every 4-6 hours when required, maximum 4000 mg/day |

» Provide supportive care, which may include the following measures.[1]

» **Oxygen**

» Prescribe oxygen if oxygen saturation <94% and maintain at target range.[1] For patients at risk of hypercapnia prescribe oxygen if oxygen saturation <88%.

- **Target oxygen saturation range of**: 
  - 94% to 96% in acutely ill patients who are not at risk of hypercapnia.[72] 
  - 88% to 92% in patients at risk of hypercapnia.[71]

- **Measure arterial blood gases** in those with SpO₂ <94%, those with a risk of hypercapnic ventilatory failure (CO₂ retention), and all patients with high-severity CAP.[1]

»

» **Fluid resuscitation**

- **Assess all patients for volume depletion** and give intravenous fluids if required.[1]

»

» **Venous thromboembolism (VTE) prophylaxis**

- Consider prophylaxis for VTE with a low molecular weight heparin for all patients who are not fully mobile.[1]

» **Airway clearance**

- Do not treat people with uncomplicated pneumonia with traditional airway clearance techniques routinely. If needed, offer these patients advice regarding expectoration of sputum.[1]

- Consider airway clearance techniques if the patient has difficulty expectorating sputum or if they have a pre-existing lung condition.[1]

» **Analgesia**

- Give simple analgesia as appropriate (e.g., for pleuritic pain).[1]
Community-acquired pneumonia (non COVID-19)

Treatment

Acute

**consider switch to pathogen-targeted antibiotic therapy**

Treatment recommended for SOME patients in selected patient group

» **Consult with a microbiologist** about appropriate pathogen-targeted antibiotic therapy.[1]

- Switch from empirical antibiotics to pathogen-targeted antibiotics as soon as specific pathogens are identified (unless there are legitimate concerns about dual-pathogen infection).[1]

» Only about one third to one quarter of patients with CAP admitted to hospital will have their pneumonia defined microbiologically.[1] Among these patients:

- Around 14% have an atypical pathogen, of which:[21]
  - 7% have *Mycoplasma pneumoniae*
  - 4% have *Chlamydophila pneumoniae*
  - 3% have *Legionella pneumophila*

- Those with infections due to *Mycoplasma*, *Chlamydophila*, and *Coxiella burnetii* will be diagnosed late in the illness on the basis of seroconversion, reducing the opportunity for early targeted therapy.[1]

» Consider switching the choice of agent once the results of sensitivity testing are available or following consultation with a microbiologist, intensivist, or respiratory physician.[1]

» **BTS recommendations for pathogen-targeted antibiotics** [1]

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Mycoplasma pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Chlamydophila pneumoniae</em></td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td><em>Legionella species</em></td>
<td>Fluoroquinolone (orally or intravenously)</td>
<td>Clarithromycin (orally or intravenously) or Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
<td>Alternative antibiotic</td>
</tr>
<tr>
<td>--------------------------------</td>
<td>-----------------------------------------------------------</td>
<td>------------------------------------------------------------</td>
</tr>
<tr>
<td><em>Streptococcus pneumoniae</em></td>
<td>Amoxicillin (orally) or benzyl/penicillin (intravenously)</td>
<td>Clarithromycin (orally) or cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
</tr>
<tr>
<td><em>Chlamydia psittaci</em></td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td><em>Coxiella burnetii</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>Haemophilus influenzae</em></td>
<td>Non-beta-lactamase-producing : amoxicillin (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td></td>
<td>Beta-lactamase-producing : amoxicillin/ clavulanate (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td><em>Gram-negative enteric bacilli</em></td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
<td>Fluoroquinolone (intravenously) or imipenem/ cilastatin (intravenously) or meropenem (intravenously)</td>
</tr>
<tr>
<td><em>Pseudomonas aeruginosa</em></td>
<td>Ceftazidime (intravenously) plus</td>
<td>Ciprofloxacin (intravenously) or piperacillin/ tazobactam (intravenously)</td>
</tr>
</tbody>
</table>
## Community-acquired pneumonia (non COVID-19)

### Treatment

#### Acute

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Gentamicin or tobramycin (dose monitoring required)</td>
<td>plus Gentamicin or tobramycin (dose monitoring required)</td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em>: non-MRSA</td>
<td>Flucloxacillin (intravenously)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>with or without</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus aureus</em>: MRSA</td>
<td>Vancomycin (intravenously; dose monitoring required)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>or linezolid (intravenously)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>or teicoplanin (intravenously)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>with or without</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
</tbody>
</table>

**confirmed CAP on chest x-ray: presenting in the community**

1st continue empirical antibiotics or switch to pathogen-targeted antibiotic therapy

» Continue empirical antibiotics in patients with CAP confirmed by chest x-ray in the community. However, where a pathogen has been identified, follow your local antibiotic protocol for the identified organism(s). [1]

» BTS recommendations for pathogen-targeted antibiotics [1]
### Acute Community-acquired pneumonia (non COVID-19) Treatment

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Preferred antibiotic</th>
<th>Alternative antibiotic</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mycoplasma pneumoniae</td>
<td>Clarithromycin (orally or intravenously)</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td>Chlamydia pneumoniae</td>
<td>Doxycycline (orally) or a fluoroquinolone (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td>Legionella species</td>
<td>Fluoroquinolone (orally or intravenously)</td>
<td>Clarithromycin (orally or intravenously) or azithromycin (in countries where it is used for managing pneumonia)</td>
</tr>
<tr>
<td>Streptococcus pneumoniae</td>
<td>Amoxicillin (orally) or benzylpenicillin (intravenously)</td>
<td>Clarithromycin (orally) or cefuroxime or cefotaxime or ceftriaxone (intravenously)</td>
</tr>
<tr>
<td>Chlamydia psittaci</td>
<td>Doxycycline (orally)</td>
<td>Clarithromycin (orally or intravenously)</td>
</tr>
<tr>
<td>Haemophilus influenzae</td>
<td>Non-beta-lactamase-producing : amoxicillin (orally or intravenously)</td>
<td>Cefuroxime or cefotaxime or ceftriaxone (intravenously) or a fluoroquinolone (orally or intravenously)</td>
</tr>
<tr>
<td>Gram-negative enteric bacilli</td>
<td>Cefuroxime or cefotaxime</td>
<td>Fluoroquinolone (intravenously) or imipenem/</td>
</tr>
<tr>
<td>Pathogen</td>
<td>Preferred antibiotic</td>
<td>Alternative antibiotic</td>
</tr>
<tr>
<td>-------------------------------</td>
<td>------------------------------------------</td>
<td>----------------------------------------------------</td>
</tr>
<tr>
<td></td>
<td>or ceftriaxone (intravenously)</td>
<td>cilastatin (intravenously) or meropenem (intravenously)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pseudomonas aeruginosa</td>
<td>Ceftazidime (intravenously)</td>
<td>Ciprofloxacin (intravenously) or piperacillin/tazobactam (intravenously)</td>
</tr>
<tr>
<td></td>
<td>plus</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Gentamicin or tobramycin (dose monitoring required)</td>
<td>Gentamicin or tobramycin (dose monitoring required)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Staphylococcus aureus: non-MRSA</td>
<td>Fluoxacillin (intravenously) with or without</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Rifampicin (orally or intravenously)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Staphylococcus aureus: MRSA</td>
<td>Vancomycin (intravenously; dose monitoring required) or linezolid (intravenously) or teicoplanin (intravenously) with or without</td>
<td>Rifampicin (orally or intravenously)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**TREATMENT**

This PDF of the BMJ Best Practice topic is based on the web version that was last updated: Jun 04, 2020. BMJ Best Practice topics are regularly updated and the most recent version of the topics can be found on bestpractice.bmj.com. Use of this content is subject to our disclaimer. © BMJ Publishing Group Ltd 2020. All rights reserved.
### Acute

In community settings, the diagnosis of CAP is based on signs and symptoms of lower respiratory tract infection, focal chest signs, and illness severity, and management is based on a suspected diagnosis. However, a chest x-ray is indicated in the community if:

- There is diagnostic doubt
- **The patient is deemed to be at risk** of underlying lung pathology (e.g., they have risk factors for lung cancer)
- Progress following treatment is not satisfactory at review.
Emerging

Coenzyme Q10

Coenzyme Q10 is a naturally occurring, fat-soluble, vitamin-like quinone that may be beneficial in people aged >60 years with CAP when given as an adjunct to antibiotic treatment. Results from a small (n=141) randomised controlled trial showed a significantly faster decline in fever, shorter hospital stays, and less treatment failure in patients with CAP receiving coenzyme Q10 (200 mg/day) compared with those receiving placebo when given for 14 days together with antibiotic therapy (prescribed according to latest guidelines).[203] Adverse events in the two groups were few and similar. CAP was diagnosed according to defined clinical and radiological criteria. It is not clear, however, whether this study had enough power to detect any meaningful differences between groups, and therefore no recommendations can be made until further evidence is available.

Omadacycline

A new modernised tetracycline antibiotic with broad-spectrum activity, designed to overcome tetracycline resistance. It is approved by the US Food and Drug Administration for the treatment of adults with CAP. It is available in oral and intravenous formulations, and is expected to be available commercially in early 2019. Like other antibiotics in the tetracycline class, omadacycline may cause discoloration of deciduous teeth, and inhibition of fetal bone growth when administered during pregnancy.

Ceftaroline

Ceftaroline is a fifth-generation parenteral extended-spectrum cephalosporin that binds to penicillin-binding proteins and prevents the synthesis of the bacterial cell wall. It has antimicrobial activity against gram-positive organisms, including Streptococcus pneumoniae, Streptococcus pyogenes, Staphylococcus aureus (including methicillin-resistant S. aureus [MRSA], vancomycin-resistant S. aureus [VRSA], and hetero-resistant vancomycin intermediate S. aureus [hVISA]), as well as many common gram-negative organisms, such as Haemophilus influenzae and Moraxella catarrhalis. Reviews have found that ceftaroline is superior to ceftriaxone in patients with CAP in terms of clinical cure rates.[204] [205]

Ceftobiprole

Ceftobiprole is a broad-spectrum parenteral cephalosporin that has microbiological activity against most typical bacterial pathogens causing CAP, including MRSA. A phase III study found that ceftobiprole was non-inferior to ceftriaxone with or without linezolid for the treatment of CAP.[206]

Nemonoxacin

A non-fluorinated, broad-spectrum quinolone. It has greater antimicrobial activity than the fluoroquinolones (e.g., levofloxacin) against MRSA, methicillin-sensitive Staphylococcus epidermidis (MSSE), methicillin-resistant S. epidermidis (MRSE), S. pneumoniae, and Enterobacter faecalis. A phase II study found that nemonoxacin has high clinical cure rates in adults with CAP.[207]

Solithromycin

A fluoroketolide with antimicrobial activity against gram-positive and gram-negative bacteria commonly associated with CAP. A completed phase II study showed that solithromycin had similar efficacy to that of levofloxacin in adults with bacterial CAP with pneumonia severity index scores of II to IV.[208] It has also been found to be non-inferior to moxifloxacin.[209] Solithromycin is currently in phase III development for the treatment of bacterial CAP.

Cethromycin
A fluoroketolide with a reported high antimicrobial activity against gram-positive and gram-negative bacteria, and atypical pathogens (including *Mycoplasma* and *Ureaplasma*). It also has in vitro activity against penicillin-resistant and macrolide-resistant gram-positive organisms, possibly due to a high affinity for the target site on the ribosomal unit.[210]
Recommendations

Monitoring

In hospital

Discuss with a senior colleague any patient who does not improve as expected. [1]

- Consider repeat chest radiograph, C-reactive protein, white cell count, and further specimens for microbiology in patients not progressing satisfactorily after 3 days of treatment.[1]
- Consider referral to a respiratory physician. [1]

Practical tip

The main reasons why patients do not improve as expected include:[1]

- Incorrect diagnosis or complicating condition (e.g., pulmonary embolism, bronchial carcinoma, bronchiectasis)
- Unexpected pathogen or pathogens not covered by antibiotic choice (e.g., ‘atypical’ pathogens, pathogens resistant to commonly used antibiotics such as ampicillin-resistant *Haemophilus influenzae*)
- Antibiotic ineffective or causing allergic reaction (e.g., poor absorption of oral antibiotic, inadequate dose, antibiotic hypersensitivity)
- Impaired local (e.g., bronchiectasis, endobronchial obstruction, aspiration) or systemic (e.g., HIV infection, myeloma) defenses
- Local (e.g., parapneumonic effusion, empyema, lung abscess) or distant (e.g., metastatic infection, septicemia, phlebitis at intravenous cannula site) complications of CAP
- Overwhelming infection
- Improvement expected too soon (e.g., in older patients).

In patients with high-severity CAP who are not responding to beta-lactam antibiotics or for whom an atypical or viral pathogen is suspected, order polymerase chain reaction (or other antigen detection test) of sputum or other respiratory tract sample.[1]

- Consider initial and follow-up viral and atypical pathogen serology.[1]

In the community

Advise patients (and their carers) to seek medical advice if their symptoms worsen rapidly or significantly; symptoms do not start to improve within 3 days; or they become systemically very unwell.[64]

- About 10% of patients managed in the community do not respond to antibiotic therapy and require hospitalisation.[122]

Admit urgently to hospital any patient on antibiotic treatment with features of moderate- or high-severity infection.[1]

Discharge and follow-up

Do not request a repeat chest radiograph before discharge from hospital in patients who have recovered satisfactorily from CAP.[1]

Arrange a follow-up visit at around 6 weeks either with the patient’s general practitioner or in a hospital clinic.[1]
Follow up

- Request a repeat chest radiograph during recovery after about 6 weeks for patients (regardless of whether they have been admitted to hospital).[1]
  - With persisting symptoms or physical signs
  - Who are at higher risk of underlying malignancy (especially smokers and those aged >50 years).
- Consider bronchoscopy in patients with persisting signs, symptoms, and radiological abnormalities at around 6 weeks after completing treatment.[1]
- Consider a chest and upper abdomen CT scan in patients with persistent signs or symptoms or with chest radiograph changes prior to bronchoscopy (e.g., if lung cancer is suspected).[98]

Patient instructions

Advise patients to rest, to drink plenty of fluids, and not to smoke.[1]

- Adequate hydration and preservation of the cough reflex during convalescence are important. The cough reflex is necessary to remove micro-organisms from the respiratory tract before it reaches the lung.
- Explain that adherence to treatment is important in patients with CAP, even after they experience clinical improvement.

At discharge, offer patients access to information about CAP, such as a patient information leaflet.[1] For all patients with CAP who smoke, offer advice according to national smoking cessation guidelines. [National Institute for Health and Care Excellence: stop smoking interventions and services]

In the community, advise patients (and their carers) to seek medical advice if their symptoms worsen rapidly or significantly; symptoms do not start to improve within 3 days; or they become systemically very unwell.[64]

- Around 10% of patients managed in the community do not respond to antibiotic therapy and require hospitalisation.[122]
## Complications

<table>
<thead>
<tr>
<th>Complications</th>
<th>Timeframe</th>
<th>Likelihood</th>
</tr>
</thead>
<tbody>
<tr>
<td>septic shock</td>
<td>short term</td>
<td>medium</td>
</tr>
<tr>
<td>Commonly complicates severe CAP. Patients have fever, leukocytosis, tachypnoea, tachycardia. Can progress rapidly to multi-organ failure and shock. It is often fatal, and survival is dependent on a high index of suspicion, early recognition, and immediate intervention.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>acute respiratory distress syndrome (ARDS)</td>
<td>short term</td>
<td>medium</td>
</tr>
<tr>
<td>Pneumonia can be complicated by ARDS, which is a condition of non-cardiogenic pulmonary oedema and severe lung inflammation. Associated with a 30% to 50% mortality, and treated with low tidal volume plateau pressure limited mechanical ventilation.[6]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>antibiotic-associated Clostridium difficile colitis</td>
<td>short term</td>
<td>medium</td>
</tr>
<tr>
<td>May occur as a result of interruption of the normal bowel flora from antibiotic use. Patients generally have diarrhoea, abdominal pain, and leukocytosis. Stool immunoassay for C difficile enzymes is diagnostic. Ideally, causative antibiotics should be stopped, and treatment is with oral metronidazole, vancomycin, or fidaxomicin.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>heart failure</td>
<td>short term</td>
<td>medium</td>
</tr>
<tr>
<td>The incidence of heart failure in hospitalised patients with CAP was 14.1% in one study.[214] There is little information about risk factors for the occurrence of cardiac complications in patients with CAP. Older age, pre-existing congestive heart failure, severity of CAP, and the use of insulin by glucose sliding scales in hospitalised patients are possible risk factors.[215] [216] [217] In patients with known cardiovascular disease, use of pneumococcal and influenza vaccine may reduce morbidity and mortality.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>acute coronary syndrome</td>
<td>short term</td>
<td>low</td>
</tr>
<tr>
<td>The incidence of acute coronary syndrome in hospitalised patients with CAP was 5.3% in one study.[214]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>cardiac arrhythmias</td>
<td>short term</td>
<td>low</td>
</tr>
<tr>
<td>The incidence of incident cardiac arrhythmia in hospitalised patients with CAP was 4.7% in one study.[214]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>necrotising pneumonia</td>
<td>short term</td>
<td>low</td>
</tr>
<tr>
<td>Regarded as a rare complication of CAP in adults. Associated with pathogens such as Staphylococcus aureus, Streptococcus pyogenes, Nocardia species, Klebsiella pneumoniae, and Streptococcus pneumoniae. Smoking, alcoholism, old age, diabetes mellitus, chronic lung diseases, or liver disease are risk factors associated with necrotising pneumonia.[221]</td>
<td></td>
<td></td>
</tr>
<tr>
<td>pleural effusion</td>
<td>variable</td>
<td>high</td>
</tr>
<tr>
<td>May occur in up to 57% of hospitalised pneumonia patients.[218] [219] About 1% to 2% of CAP cases with pleural effusion are complicated with empyema.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
### Complications

<table>
<thead>
<tr>
<th>Complication</th>
<th>Timeframe</th>
<th>Likelihood</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pleural effusion</td>
<td>variable</td>
<td>low</td>
</tr>
<tr>
<td>Lung abscess</td>
<td>variable</td>
<td>low</td>
</tr>
<tr>
<td>Pneumothorax</td>
<td>variable</td>
<td>low</td>
</tr>
</tbody>
</table>

Pleural effusion is considered to be an indicator of pneumonia severity and is clearly associated with an increased risk of treatment failure.\[5\] [1] [220]

A rare complication, frequently requiring prolonged antibiotic therapy and, in some cases, surgical drainage.

A rare complication of CAP in adults. Pneumothorax is associated with bacterial pneumonia caused by staphylococcus, streptococcus, and other type of bacteria, which may cause the collapse of a lung.

### Prognosis

For patients admitted to hospital, mortality rate ranges from 5% to 15%, but increases to 20% to 50% in patients requiring admission to the intensive care unit (ICU).\[6\] [106] Patients treated in the community generally have a good prognosis.\[1\]

Risk factors associated with increased 30-day mortality include bacteremia, admission to the ICU, comorbidities (especially neurological disease), and infection with a potentially multidrug-resistant pathogen (e.g., *Staphylococcus aureus*, *Pseudomonas aeruginosa*, Enterobacteriaceae).\[36\] [123] [124] [125]

Readmission rates in patients with CAP range from 7% to 12%.\[211\] [212] In most cases, exacerbation of comorbidities (mainly cardiovascular, pulmonary, or neurological disease) is responsible for readmission.

Prognostic biomarkers such as pro-adrenomedullin, prohormone forms of atrial natriuretic peptide, cortisol, procalcitonin, and C-reactive protein are being studied as predictors of mortality; however, further studies are required before these biomarkers are used for this function in clinical practice.\[213\]
## Diagnostic guidelines

### Europe

**Pneumonia in adults: diagnosis and management (withdrawn during COVID-19 pandemic)**
- **Published by:** National Institute for Health and Care Excellence
- **Last published:** 2019

**BTS guideline for diagnostic flexible bronchoscopy in adults**
- **Published by:** British Thoracic Society
- **Last published:** 2013

**BTS guidelines for the management of community acquired pneumonia in adults**
- **Published by:** British Thoracic Society
- **Last published:** 2009

### North America

**Diagnosis and treatment of adults with community-acquired pneumonia**
- **Published by:** American Thoracic Society; Infectious Diseases Society of America
- **Last published:** 2019

### Asia

**Diagnosis and treatment of community-acquired pneumonia in adults**
- **Published by:** Chinese Thoracic Society; Chinese Medical Association
- **Last published:** 2017

### Oceania

**Community acquired pneumonia**
- **Published by:** The Royal Children’s Hospital Melbourne
- **Last published:** 2016
# Treatment guidelines

## Europe

<table>
<thead>
<tr>
<th>Topic</th>
<th>Published by</th>
<th>Last published</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pneumonia (community-acquired): antimicrobial prescribing</td>
<td>National Institute for Health and Care Excellence</td>
<td>2019</td>
</tr>
<tr>
<td>Guidelines for the management of adult lower respiratory tract infections</td>
<td>European Respiratory Society</td>
<td>2011</td>
</tr>
<tr>
<td>BTS guidelines for the management of community acquired pneumonia in adults</td>
<td>British Thoracic Society</td>
<td>2009</td>
</tr>
</tbody>
</table>

## North America

<table>
<thead>
<tr>
<th>Topic</th>
<th>Published by</th>
<th>Last published</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diagnosis and treatment of adults with community-acquired pneumonia</td>
<td>American Thoracic Society; Infectious Diseases Society of America</td>
<td>2019</td>
</tr>
</tbody>
</table>
## Asia

**Guideline for antibiotic use in adults with community-acquired pneumonia**

*Published by:* Korean Society of Infectious Diseases; Korean Society for Chemotherapy  *Last published:* 2018

**Diagnosis and treatment of community-acquired pneumonia in adults**

*Published by:* Chinese Thoracic Society; Chinese Medical Association  *Last published:* 2017

**Executive summary of the Gulf Cooperation Council practice guidelines for the management of community-acquired pneumonia**

*Published by:* Gulf Cooperation Council Community-acquired Pneumonia Working Group  *Last published:* 2007

**Guidelines for the management of community acquired pneumonia in adults, revised edition**

*Published by:* Committee for The Japanese Respiratory Society  *Last published:* 2006

## Africa

**Guideline for management of community-acquired pneumonia in adults**

*Published by:* South African Thoracic Society  *Last published:* 2017

## Oceania

**Community acquired pneumonia**

*Published by:* The Royal Children’s Hospital Melbourne  *Last published:* 2016
Online resources


2. Public Health England: complete routine immunisation schedule (external link)

3. National Institute for Health and Care Excellence: stop smoking interventions and services (external link)

4. Abbreviated Mental Test Score (external link)
Key articles


References


Community-acquired pneumonia (non COVID-19)


Community-acquired pneumonia (non COVID-19)


Community-acquired pneumonia (non COVID-19)

References


68. Royal College of Physicians. National early warning score (NEWS) 2: standardising the assessment of acute-illness severity in the NHS. Dec 2017 [internet publication]. Full text


70. Surviving Sepsis Campaign. Hour-1 bundle. 2018 [internet publication]. Full text


Community-acquired pneumonia (non COVID-19)


Community-acquired pneumonia (non COVID-19)

criteria be used in older people? A compilation study of two prospective cohorts. Age Ageing. 2006 May;35(3):286-91. Full text Abstract


132. Fritz Z, Slowther AM, Perkins GD. Resuscitation policy should focus on the patient, not the decision. BMJ. 2017 Feb 28;356:j813 Full text Abstract


<table>
<thead>
<tr>
<th>Reference</th>
<th>Text</th>
</tr>
</thead>
<tbody>
<tr>
<td>150.</td>
<td>Pendlebury ST, Klaus SP, Mather M, et al. Routine cognitive screening in older patients admitted to acute medicine: abbreviated mental test score (AMTS) and subjective memory complaint versus Montreal Cognitive Assessment and IQCODE. Age Ageing. 2015 Oct 13;44(6):1000-5 Full text Abstract</td>
</tr>
</tbody>
</table>


155. Nova Scotia Health Authority. This is not my Mom. 2012 [internet publication] Full text


Community-acquired pneumonia (non COVID-19)


190. Flowers L. Nicotine replacement therapy. Am J Psychiatry Resid. 2016 Jun;11(6)4-7  Full text


193. National Centre for smoking cessation and training. Smoking cessation and mental health. 2014 [internet publication].  Full text


Community-acquired pneumonia (non COVID-19)

References


Images

Figure 1: Posterior-anterior chest radiograph showing right upper lobe consolidation in a patient with community-acquired pneumonia

Figure 2: Chest radiograph showing left upper lobe cavitating pneumonia

From the collection of Dr Jonathan Bennett. Used with permission
Figure 3: Left-sided pleural effusion

From the collection of Dr R Light. Used with permission
Figure 4: Increased opacification of the right perihilar region and superior segment of the right lower and upper lobes consistent with worsening aspiration pneumonia

From the collection of Dr Roy Hammond. Used with permission
Figure 5: Risk assessment and management of CAP in the first 4 hours: hospital setting

Community setting: manage according to clinical judgement + CRB-65 score

0 Low-severity
- Likely suitable for home treatment*
  - Oral amoxicillin preferred
    - plus rest, smoking cessation, fluids, and analgesics if pain

1 - 2 Moderate-severity
- Consider hospital referral**
  - Do not give antibiotics prior to hospital referral

3 - 4 High-severity
- Urgent hospital admission
- Prior to transfer: Consider giving intravenous benzylpenicillin or oral amoxicillin if life-threatening or if a delay in hospital treatment of >6 hours is likely

---

Figure 6: Risk assessment and management of CAP in the first 4 hours: community setting. *Consider social circumstances, unstable comorbidities, ability to take oral treatment safely and reliably, and patient’s wishes when deciding whether to treat patients at home. **The majority of patients with moderate-severity CAP should be referred to hospital. However, you should consider social circumstances, ability to take oral treatment safely, unstable comorbidities, and patient’s wishes when deciding on home treatment#

Disclaimer

BMJ Best Practice is intended for licensed medical professionals. BMJ Publishing Group Ltd (BMJ) does not advocate or endorse the use of any drug or therapy contained within this publication nor does it diagnose patients. As a medical professional you retain full responsibility for the care and treatment of your patients and you should use your own clinical judgement and expertise when using this product.

This content is not intended to cover all possible diagnosis methods, treatments, follow up, drugs and any contraindications or side effects. In addition, since such standards and practices in medicine change as new data become available, you should consult a variety of sources. We strongly recommend that you independently verify specified diagnosis, treatments and follow-up and ensure it is appropriate for your patient within your region. In addition, with respect to prescription medication, you are advised to check the product information sheet accompanying each drug to verify conditions of use and identify any changes in dosage schedule or contraindications, particularly if the drug to be administered is new, infrequently used, or has a narrow therapeutic range. You must always check that drugs referenced are licensed for the specified use and at the specified doses in your region.

Information included in BMJ Best Practice is provided on an “as is” basis without any representations, conditions or warranties that it is accurate and up to date. BMJ and its licensors and licensees assume no responsibility for any aspect of treatment administered to any patients with the aid of this information. To the fullest extent permitted by law, BMJ and its licensors and licensees shall not incur any liability, including without limitation, liability for damages, arising from the content. All conditions, warranties and other terms which might otherwise be implied by the law including, without limitation, the warranties of satisfactory quality, fitness for a particular purpose, use of reasonable care and skill and non-infringement of proprietary rights are excluded.

Figure 1 – BMJ Best Practice Numeral Style

5-digit numerals: 10,000
4-digit numerals: 1000
numerals < 1: 0.25

Where BMJ Best Practice has been translated into a language other than English, BMJ does not warrant the accuracy and reliability of the translations or the content provided by third parties (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages). BMJ is not responsible for any errors and omissions arising from translation and adaptation or otherwise. Where BMJ Best Practice lists drug names, it does so by recommended International Nonproprietary Names (rINNs) only. It is possible that certain drug formularies might refer to the same drugs using different names.

This approach is in line with the guidance of the International Bureau of Weights and Measures Service.

Contact us
+ 44 (0) 207 111 1105
support@bmj.com

BMJ
BMA House
Tavistock Square
London
WC1H 9JR
UK
Contributors:

// Peer Reviewers:

Wei Shen Lim,
Consultant Respiratory Physician and Honorary Professor of Medicine
Nottingham University Hospitals NHS Trust, Nottingham, UK
DISCLOSURES: WSL was chairman of the British Thoracic Society community-acquired pneumonia guidelines committee and a member of the guideline development group for the National Institute for Health and Care Excellence pneumonia guidelines. His institution has received unrestricted investigator-initiated research funding from Pfizer for a multicentre study of pneumococcal pneumonia in which he was the chief investigator, and research funding from the National Institute for Health Research for studies in pneumonia in which he was the principal investigator. He is also an author of at least one reference cited in the topic.

// Expert Advisers:

Jonathan Bennett, MD
Honorary Professor of Respiratory Sciences
University of Leicester, Respiratory Consultant, Glenfield Hospital, Leicester, UK
DISCLOSURES: JB declares that he has no competing interests.

Acknowledgements,

BMJ Best Practice would like to gratefully acknowledge the previous team of expert contributors, whose work has been retained in parts of the content:
Catia Cilloniz MSc, PhD, Post-doctoral Research, Pneumology Department, Hospital Clinic of Barcelona, CIBERES, IDIBAPS, Barcelona, Spain, Antoni Torres MD, PhD, Professor of Medicine, Director, Pulmonary Intensive Care Unit, Respiratory Institute, Hospital Clinic of Barcelona, Barcelona, Spain
DISCLOSURES: CC and AT are each authors of a number of references cited in this topic.

// Editors:

Helena Delgado-Cohen,
Section Editor, BMJ Best Practice
DISCLOSURES: HDC declares that she has no competing interests.

Rachel Wheeler,
Lead Section Editor, BMJ Best Practice
DISCLOSURES: RW declares that she has no competing interests.

Julie Costello,
Comorbidities Editor, BMJ Best Practice
DISCLOSURES: JC declares that she has no competing interests.

Adam Mitchell,
Drug Editor, BMJ Best Practice
DISCLOSURES: AM declares that he has no competing interests.