PNEUMONIA PART 1

DR UMMAR

Pneumonia

An acute respiratory illness associated with recently

developed radiological pulmonary shadowing which

may be segmental, lobar or multilobar.

Or,

Inflammation in the lung characterized by accumulation

of secretions and inflammatory cells in alveoli.



CLASSIFICATION

Pneumonia: Classifications

Clinically

Community-acquired pneumonia (CAP): Onset in community or during 1st 2 days of hospitalization (Strep. pneumoniae most common)

Hospital-acquired Pneumonia(HAP/nosocomial): Occurring 48 hrs after hospitalization

Suppurative & Aspiration pneumonia

Pneumonia in immunocompromised patient: caused by opportunistic organisms (Pneumocystis jirovecii).

Pneumonia: Classifications..

Anatomically

Lobar pneumonia if one or more lobe is involved

Broncho-pneumonia (Lobular)

1.more patchy alveolar consolidation associated with bronchial and bronchiolar inflammation often affecting both lower lobes

2.the pneumonic process has originated in one or more bronchi and extends to the surrounding lung tissue

Pneumonia: Classifications..

According to causes

- Bacterial (the most common cause of pneumonia)
- ➢ Viral pneumonia
- ➤Fungal pneumonia
- ➤Aspiration pneumonia
- >Chemical pneumonia (ingestion of kerosene or inhalation of

irritating substance)

Pneumonia: Classifications..

>Typical pneumonia:

Respiratory symptoms are more than constitutional symptoms > Atypical pneumonia: Constitutional symptoms are more than respiratory symptoms

(Behaviourist's classification)

Easy pneumonia (responds to initial treatment)

Difficult pneumonia (fails to do so)

COMMUNITY-ACQUIRED PNEUMONIA (CAP)

Community-acquired pneumonia (CAP)

- >Onset in community or during 1st 2 days of hospitalization
- ➢ Strep. pneumoniae most common 50%
- ➢ It affects all age groups but is particularly common at the extremes of age.
- >Worldwide, CAP continues to kill more children than any other
- illness, and its propensity to ease the passing of the frail and
- elderly led to pneumonia being known as the 'old man's friend'.

Community-acquired pneumonia (CAP)..

≻Most cases are spread by droplet infection.

≻ May occur in previously healthy individuals.

Streptococcus pneumoniae remains the most common infecting agent.

>Other organisms may be involved which depends on the age of the

patient and the clinical context.

 \succ Viral infections are important causes of CAP in children, and their

contribution to adult CAP is increasingly recognized

Community-acquired pneumonia (CAP)..

Mycoplasma pneumoniae is more common in young people and rare in the elderly.

Haemophilus influenzae is more common in the elderly, particularly when underlying lung disease is present.

Legionella pneumophila occurs in local outbreaks centred on contaminated cooling towers in hotels, hospitals and other industrial buildings.

Staphylococcus aureus is more common following an episode of influenza. Community-acquired pneumonia (CAP)..

Factors that predispose to pneumonia

- Cigarette smoking
- Upper respiratory tract infections
- ≻ Alcohol
- Corticosteroid therapy
- ≻Old age

- Recent influenza infection
- Pre-existing lung disease
 HIV
 Indoor air pollution

Community-acquired pneumonia (CAP)...

Organisms causing CAP

Bacteria

- Streptococcus pneumoniae Coxiella burnetii
- *Mycoplasma pneumoniae* (Q fever, 'querry' fever)
- Legionella pneumophila
- Chlamydia pneumoniae
- Haemophilus influenzae
- Staphylococcus aureus
- Chlamydia psittaci

- Klebsiella pneumoniae
 - (Freidländer's bacilus)
- Actinomyces israelii

- Viruses
- > Influenza, parainfluenza
- > Measles
- > Herpes simplex
- > Varicella
- > Adenovirus
- > Cytomegalovirus (CMV)
- Coronavirus (Urbani SARSassociated coronavirus)



PATHOPHYSIOLOGY

Pneumonia: mode of transmission

> Bacteria and viruses living in your nose, sinuses, or mouth may spread to your lungs

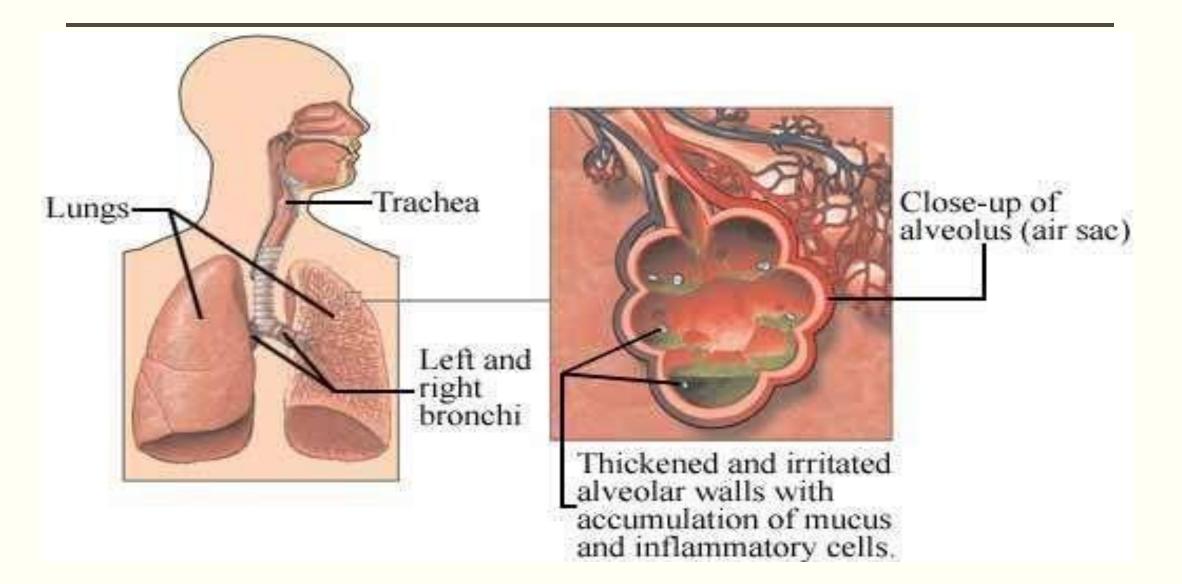
- You may breathe some of these germs directly into your lungs (droplets infection)
- You breathe in (inhale) food, liquids, vomit, or fluids from the mouth into your lungs (aspiration pneumonia)

- The streptococci reach the alveoli and lead to inflammation and pouring of an exudates into the air spaces
- WBCs migrates to alveoli, the alveoli become more thick due to its filling consolidation, involved areas by inflammation are not adequately ventilated, due to secretion and edema
- This will lead to partial occlusion of alveoli and bronchi causing a decrease in alveolar oxygen content

Venous blood that goes to affected areas without being oxygenated and returns to the heart (ventilation-perfusion mismatch)

➢This will lead to arterial hypoxemia and even death due to interference with ventilation

Pathophysiology..



CLINICAL FEATURES

Pneumonia, particularly lobar pneumonia, usually presents as an acute illness.

Systemic features such as fever, rigors, shivering and malaise predominate and delirium may be present.

>The appetite is invariably lost and headache frequently reported.

>Pulmonary symptoms include cough, which at first is

characteristically short, painful and dry, but later accompanied by the expectoration of mucopurulent sputum.

Rust-coloured sputum may be seen in patients with Strep. pneumoniae, and the occasional individual may report haemoptysis.

Pleuritic chest pain may be a presenting feature and, on occasion, may be referred to the shoulder or anterior abdominal

> Upper abdominal tenderness is sometimes apparent in patients

with lower lobe pneumonia or if there is associated hepatitis.

Less typical presentations may be seen in the very young and the elderly.

On examination,

The respiratory and pulse rate may be raised and the blood pressure low, while an assessment of the mental state may reveal a delirium. These are important indicators of the severity of the illness

Not all patients are pyrexial but this is a helpful diagnostic clue if present.

Oxygen saturation on air may be low, and the patient cyanosed and distressed.

On examination..

> Chest signs vary, depending on the phase of the inflammatory response.

> When consolidated, the lung is typically dull to percussion and, as conduction of sound is enhanced, auscultation reveals bronchial breathing and whispering pectoriloquy; crackles are heard throughout.

However, in many patients, signs are more subtle with reduced air entry only, but crackles are usually present.

On examination..

> An assessment of nutrition is important as, if poor, the response to treatment will be impaired, particularly in the elderly.

> On occasion, inferences as to the likely organism may be drawn from clinical examination. For example, the presence of herpes labialis may point to streptococcal infection, as may the finding of 'rusty' sputum.

> The presence of poor dental hygiene should prompt consideration of Klebsiella or Actinomyces israelii.

Chronic Pneumonia

- > Symptoms creep in slowly
- > Fever that lasts a week
- > Coughing for three weeks
- > Enlarged cervical & axillary lymphnodes

> Haemoptysis

> Recurrence of symptoms after finishing antibiotic course

Differential diagnosis of pneumonia

- > Pulmonary infarction
- > Pulmonary/pleural TB
- > Pulmonary oedema (can be unilateral)
- > Pulmonary eosinophilia
- > Malignancy: bronchoalveolar cell carcinoma
- Rare disorders: cryptogenic organising pneumonia/
- bronchiolitis obliterans organising pneumonia (COP/BOOP)
- > Venous thromboembolism, Pulmonary haemorrhage
- > ARDS
- > Drug toxicity



INVESTIGATIONS

The aims of investigation are

- > Confirm the diagnosis
- > Exclude other conditions
- > Assess the severity
- > Identify the development of complications

Full blood count

- Very high (> 20 × 109/L) or low (< 4 × 109/L) white cell count: marker of severity
- Neutrophil leucocytosis > 15 × 109/L: suggests bacterial aetiology
- Haemolytic anaemia: occasional complication of Mycoplasma
- Erythrocyte sedimentation rate/C-reactive protein: Nonspecifically elevated

> Blood culture: Bacteraemia: marker of severity

> Urea and electrolytes:

•Urea > 7 mmol/L (~20 mg/dL): marker of severity

Hyponatraemia: marker of severity

- Liver function tests:
 - •Abnormal if basal pneumonia inflames liver
 - Hypoalbuminaemia: marker of severity

 Serology: Acute and convalescent titres for Mycoplasma, Chlamydia, Legionella and viral infections
 Cold agglutinins: Positive in 50% of patients with Mycoplasma
 Arterial blood gases: Measure when SaO2 < 93% or when severe clinical features to assess ventilatory failure or acidosis

Sputum

> Sputum samples

Gram stain, culture and antimicrobial sensitivity testing. Gram stain of sputum showing Grampositive diplococci characteristic of Strep. pneumoniae.

>Oropharynx swab

PCR for *Mycoplasma pneumoniae and other atypical* **pathogens**

≻Urine

Pneumococcal and/or Legionella antigen > Pleural fluid

Always aspirate and culture when present in more than trivial amounts, preferably with ultrasound guidance

Other markers of severity of Pneumonia

- >CXR > One lobe involved
- ≻ Pao2 <8kPa
- > Low albumin(<35gm/L)</pre>
- >WBC(<4000/cmm or >20000/cmm)
- > Blood culture positive

Chest X-ray

Lobar pneumonia

- Patchy opacification evolves into homogeneous consolidation of affected lobe
- Air bronchogram (air-filled bronchi appear lucent against consolidated lung tissue) may be present.
- Bronchopneumonia: Typically patchy and segmental shadowing
 Complications: Para-pneumonic effusion, intrapulmonary
 abscess or empyema
- Staph. aureus: Suggested by multilobar shadowing, cavitation, pneumatocoeles and abscesses



For evaluation of PSI > CBC HCT, TC, DC > RBS > Blood Urea > Serum electrolytes > CXR > ABG Analysis > Pulse oximetry



MANAGEMENT

Management

The principles of management focusing on

- > Adequate oxygenation
- > Appropriate fluid balance
- > Antibiotics
- In severe or prolonged illness,
- > Nutritional support may be required

Evaluate the effectiveness of administered medications
 Explain all procedures to the patient and family

Oxygen

>Oxygen should be administered to all patients with

- tachypnoea,
- hypoxaemia,
- •hypotension or
- acidosis

The aim of maintaining the PaO2 at or above 8 kPa (60 mmHg) or the SaO2 at or above 92%.

Oxygen

 High concentrations (35% or more), preferably humidified, should be used in all patients who do not have hypercapnia associated with COPD.

•Continuous positive airway pressure (CPAP) should be considered in those who remain hypoxic despite this and these patients should be managed in a highdependency or intensive care environment, where mechanical ventilation can be rapidly employed.

Intravenous fluids

- These should be considered in patients with severe illness, older patients and those who are vomiting.
- Otherwise, an adequate oral intake of fluid should be encouraged.
- Inotropic support may be required in patients with shock

Antibiotics

Prompt administration of antibiotics improves the outcome.

Th

- e initial choice of antibiotic is guided by
- > clinical context,
- > severity assessment,
- > local knowledge of antibiotic resistance patterns
- \succ any available epidemiological information.
- The choice of empirical antibiotic therapy is considerably more challenging, due to
- Diversity of pathogens
- > Drug resistance.

Antibiotics : Uncomplicated CAP:

Amoxicilin 500 mg 3 times daily orally

> If patient is allergic to penicillin: Clarithromycin 500 mg twice daily orally

or Erythromycin 500 mg 4 times daily orally

>If Staphylococcus is cultured or suspected: Flucloxacillin 1–2 g 4 times

daily N plus Clarithromycin 500 mg twice daily N

>If Mycoplasma or Legionella is suspected: Clarithromycin 500 mg twice

daily orally or Nor Erythromycin 500 mg 4 times daily orally N plus

Rifampicin 600 mg twice daily IV in severe cases

Antibiotics : Severe CAP:

Clarithromycin 500 mg twice daily Nor Erythromycin 500 mg 4 times daily N

plus

Co-amoxiclav 1.2 g 3 times daily Nor

Ceftriaxone 1-2 g daily Nor

Cefuroxime 1.5 g 3 times daily Nor

Amoxicilin 1 g 4 times daily N plus fuctoxacil in 2 g 4 times daily N

Antibiotics: Oral antibiotics are usually adequate unless the patient has a

- > severe illness,
- > impaired consciousness,
- > loss of swallowing reflex, or
- > functional or anatomical reasons for malabsorption.

In most patients with uncomplicated pneumonia, a 7-day course is adequate, although treatment is usually required for longer in those with Legionella, staphylococcal or Klebsiella pneumonia.

Antibiotics:

- **Duration of therapy**
- 5-7 days outpatients
- 10-14 days Mycoplasma, Chlamydia, Legionella
- 14+ days chronic steroid users
- 14-21 days Staph. aureas, Legionella spp
- [Am J Respir Crit Care Med 163:1730-54, 2001]

Pain

- It is important to relieve pleural pain, as it may prevent the patient from breathing normally and coughing efficiently.
- For the majority, simple analgesia with paracetamol, co-codamol or NSAIDs is sufficient.
- In some patients, opiates may be required but these must be used with extreme caution in patients with poor respiratory function, as they may suppress ventilation.

Physiotherapy

May help expectoration in those who suppress cough because of pleural pain.

- > Maintain a patent airway and adequate oxygenation
- > Use suction if the patient can't produce a specimen
- Provide a high calorie, high protein diet & soft foods
- > Provide a quiet, calm environment, with frequent rest periods
- > Monitor the patient's ABG levels, especially if he's hypoxic
- > Assess the patient's respiratory status
- > Auscultate breath sounds at least every 4 hours

Delayed resolution means

> Physical signs persist for more than 2 weeks and

- Radiological features persist for more than 4 weeks after antibiotic therapy.
- **Non-resolution means**
- > If radiological opacity persists after 8 weeks (with treatment/after antibiotic therapy).

Delayed resolution suggests

- L the diagnosis is incorrect
- IL Incorrect microbiological diagnosis
- III. Fungal, tubercular or atypical pneumonia
- **IV. recurrent aspiration**
- V. Improper antibiotic or insufficient dose
- VI. pneumonia may be secondary to a proximal bronchial obstruction

VII. complication has occurred (Empyema or atelectasis)

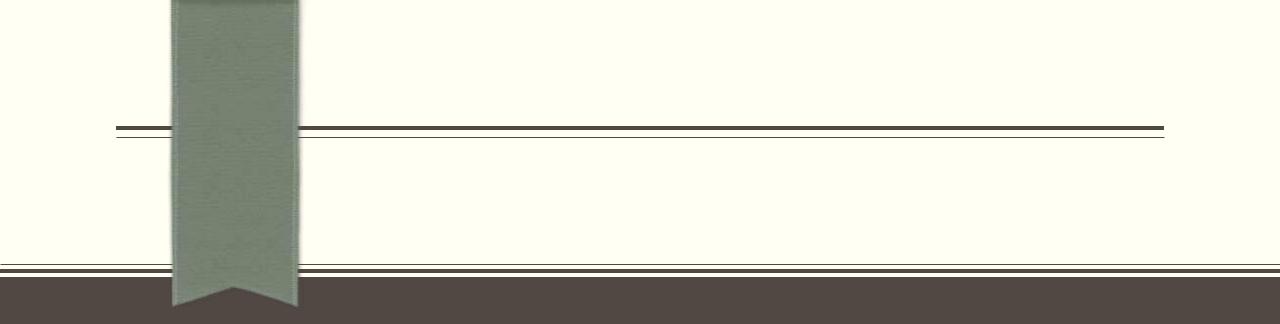
VIIL Bronchial obstruction (bronchial carcinoma, adenoma, foreign body)

IX. Immunocompromised patient (HIV, DM, lymphoma, leukemia, multiple myeloma).

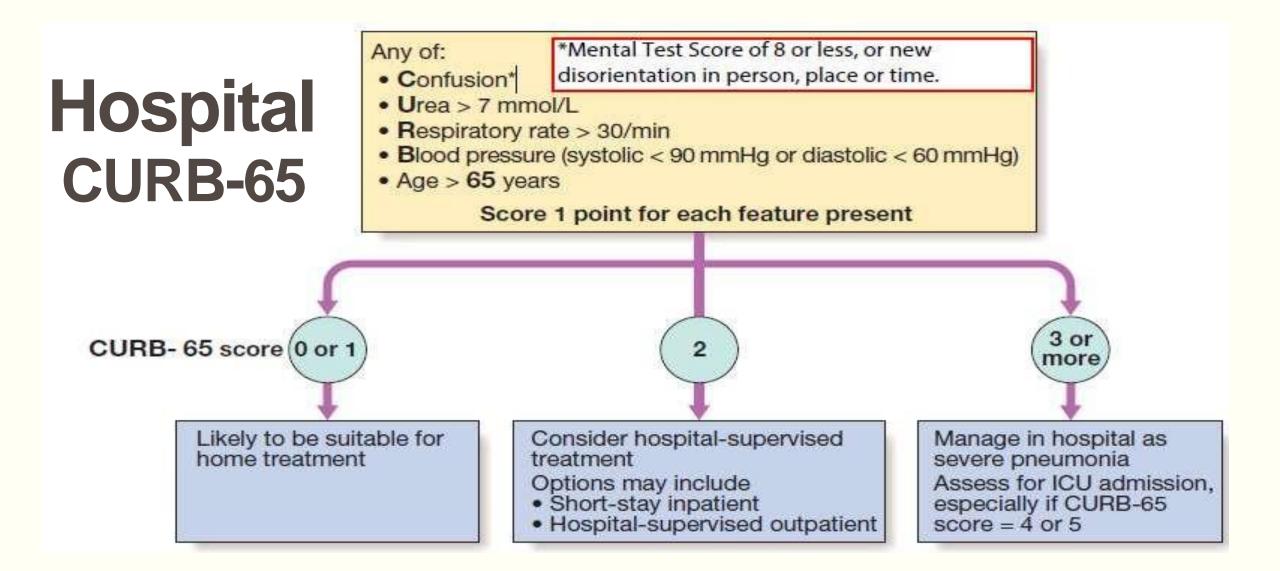
Assessment of Nonresponders

Wrong Organism Drug-resistant Pathogen: (bacteria, mycobacteria, virus, fungus Inadequate Antimicrobial Therapy Wrong Diagnosis Atelectasis Pulmonary Embolus ARDS Pulmonary Hemorrhage Underlying Disease Neoplasm

Complication Empyema or Lung Abscess Clostridium difficile Colitis Occult Infection Drug Fever



REFERRAL TO ITU



Indications for referral to ITU

- > CURB score of 4-5, failing to respond rapidly to initial
- > management
- Persisting hypoxia (PaO2 < 8 kPa (60 mmHg)),</p>
- despite high concentrations of oxygen
- > Progressive hypercaphia
- > Severe acidosis
- > Circulatory shock
- > Reduced conscious level

