Internal medicine

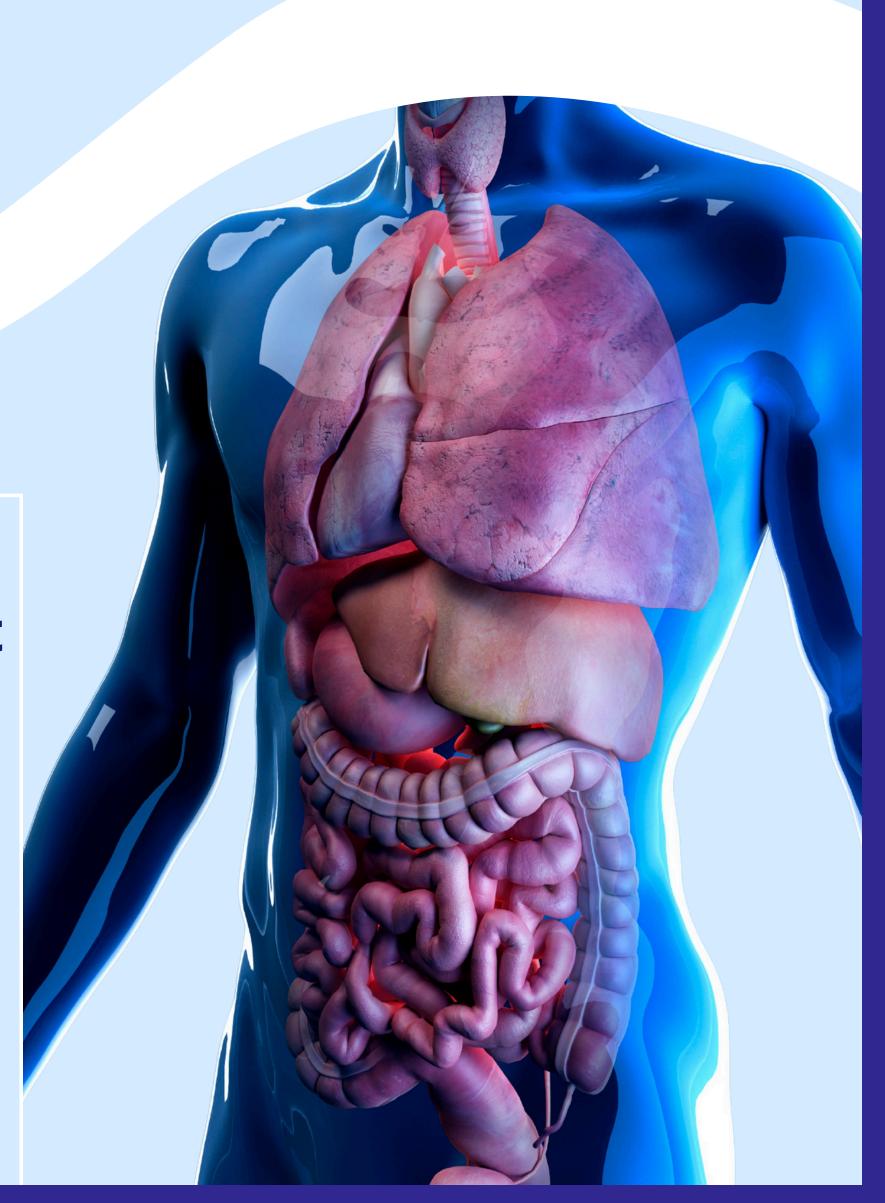


PNEUMONIA

Dr.Maha AlSadik

Done by: Aman Abu Sakout

Corrected by: Aya Abu Samra



PNEUMONIA

Definition:

It is a syndrome of acute infection of the lung parenchyma, characterized by clinical and / or radiological picture of consolidation. Commonly due to bacterial infection when the

cause is non Infectious, it is termed pneumonitis.

Classification of Pneumonia

1) Anatomical Classification:

- ☐ Lobar pneumonia.

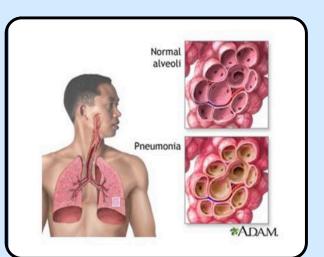
 Every lobe devide into segments (10 in RtL& 8-9 in LtL)

 ☐ Segmental or subsegmental pneumonia.
- ☐ Bronchopneumonia. Diffused and patchy
- 2) Aetiological Classification:

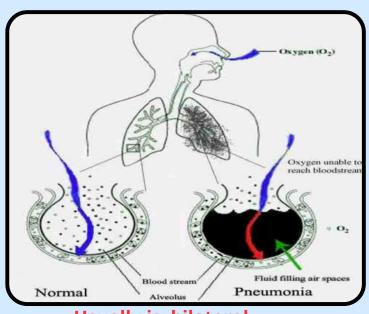
According to responsible micro organisms.

3) Environmental Classification:

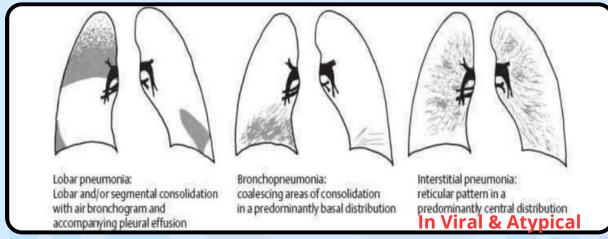
Community or hospital acquired pneumonia



Exudate will interfere gas exchanging result in complication



Usually is bilateral



pneumonia

1 Anatomical Classification:

☐ Lobar pneumonia: One or more lobes are uniformly affected by inflammation and consolidation.

☐ Segmental or subsegmental pneumonia: There is only part of the lobe is affected.

☐ Bronchopneumonia: There is a patchy involvement of lung parenchyma, particularly in lower zone

2 - Aetiological Classification:

• Bacterial:

Specific:TB

Non-specific:

- -Gram +ve organism / Gram-ve organism
- Atypical e.g. Mycoplasma and Legionella species Chlamydia,
- Viral: e.g. H1N1
- Fungal: e.g. Histoplasmosis and Aspergillosis
- Parasitic: e.g. Malaria Inflammation w/o infection
- Other causes of pneumonia:

Allergic pneumonitis: e.g., Lupus pneumonitis. May be sever need corticosteroids to treat

Chemical pneumonitis e.g.: Lipoid pneumonia. Lipid deposits in alveoli result from Oxygenous source as some people

Radiation: e.g. Radiation Pneumonitis

use oily nasal drops—> accidental inhalation then enter the lung causing chemical reactions which called lipoid pneumonia, OR may the source from endogenous (lipid metabolism)

Specially in chest radiation and will cause complications like ILF

3- Community or hospital acquired pneumonia:

Community acquired pneumonia:

> Pneumonia which is acquired in the community or at hospitalization within the first 2 days

> The most common organisms are Streptococcal pneumonia, Atypical, Staph. Aureus and Hemophilus influenza

Hospital acquired pneumonia: or Nosocomial pneumonia:

- ➤It is a pneumonia which is acquired in the hospital after 2 days of hospitalization.
- ➤The commonest organisms are G-ve bacilli e.g: pseudomonas aeruginosa, Klebsiella

and proteus

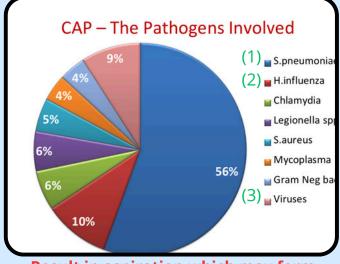
Community Acquired Pneumonia (CAF Epidemiology

- 6th leading cause of death
- 1 -25 cases / 1000 inhabitants / year
- 20% require admission
- 14% Average mortality rate
- Mortality disproportionately high in old age

Community Acquired Pneumonia (CAP)

Pneumonias – Classification.... CAP • Community Acquired HAP • Hospital Acquired VAP • Ventilator Acquired

VAP is sub of HAP but the difference is VAP come from ventilation tools after 48 hs



Result in aspiration which may form environment for growth bacteria or cause pneumonia by itself

Predisposing factors:

- 1. Impaired cough reflex e.g. Anesthesia, Alcoholism, Tracheostomy. Due to loss of consciousness
- 2. Impairment of mucociliary activity surface of cilia has a mucus, cilia Move upward to coughing up any organisms may impairment if mucus becomes thick as in cystic fibrosis Or movement is weak as in mortality cilia syndrome
- 3. Decrease of effective phagocytic activity of alveolar macrophages and neutrophils.

 Community Acquired Pneumonia (CAP)

Mode of infection and causative organisms:

A- Aspiration: Organisms in the content of aspiration

Predisposed by impaired cough:

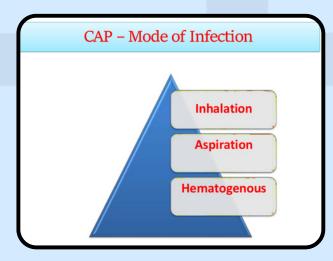
Anesthesia, Alcoholism, Tracheostomy.

B-Inhalation:

Patient to patient by direct contact through droplet infection 0r Airborne infection.

C- Colonization: In chronically ill patients e.g. COPD, Bronchiectasis.

D- Blood spread: IV. Cannula, CV lines, and IV drug abusers



Pathology of Pneumonia

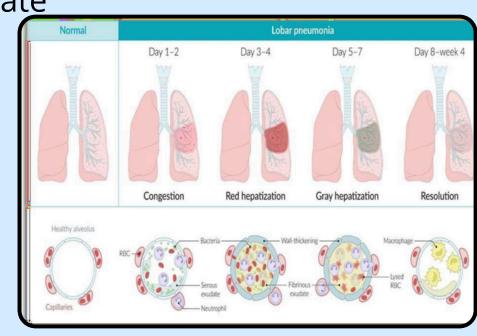
- > The commonest feature of pathology is the presence of cellular exudate in the alveolar spaces.
- > In pneumococcal and viral pneumoniae, resolution occurs through the action of macrophages and lung tissue may return to former state

Diagnosis of pneumonia

History and examination

- Symptoms:
- > Systemic: Fever, malaise, anorexia, body pain, sweating
- Chest: Cough, purulent expectoration, sometimes blood tinged, Dyspnea and Pleuritic chest pain.

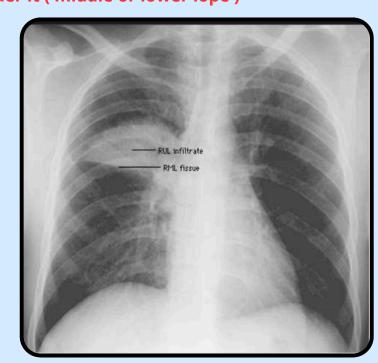
More with lower= classic pneumonia



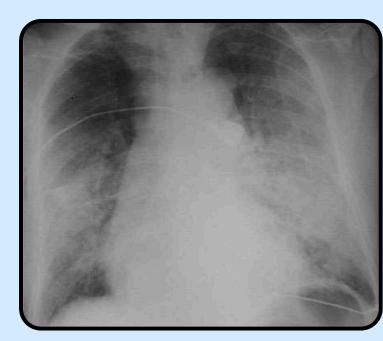
- Bacillus enter the alveoli where macrophages neutrophil are found with serous exudate
- Fluidy(serous Exudate —>fibrous exudate (more thick) + ^ in Cells
- As Red hepatization but cells breaked down
 turn to gray color
- Macrophage start lung clearance and lysis all dead tissue and microbs

Examination **CAP – Two Types of Presentations** > General: ☐ Fever **Atypical** Classical ☐ Tachypnea: Short and rapid breathing with fever Sudden onset Insidious onset ☐ Tachycardia: Relative Bradycardia High fever, chills Low grade fever, Confusion in viral pneumonia. when there is fever but no ^ in Rate as viral inf. Pleuritic chest pain, SOB **SOB** Productive cough, Rusty · Dry cough. ☐ Cyanosis in severe pneumonia. sputum, blood tinge Poor general condition Diarrhea, abdominal pain > Local chest symptoms: = Clinical picture of consolidation High mortality up to 20% • Low mortality 1-2%; except (very important) ☐ Increased TVF, in patients with in cases of Legionellosis bacteremia Mycoplasma, Chlamydia, ☐ Impaired note or Dullness to percussion, S.pneumoniae causative Legionella, and Viruses ☐ Bronchial breath sounds, or crackle bronchophony: +ve, as TVF but in auscultation **Egophony: patient say (A) we here it (E) Streptococcus Pneumonia** Most common cause of CAP (about 2/3 of cases of CAP) These are gram positive diplococci Typical presentation (e.g. fever, chills, Pleuritic chest pain, cough with rusty sputum) Lobar infiltrate on CXR **Atypical Pneumonia** > Legionella pneumonia ☐ Legionnaires' disease is a lung infection occur by inhaling the bacteria from contaminated water system like air conditioning or hot tubs. □ Older adults, smokers and people with weakened immune systems are particularly susceptible. On X-Ray = interstitial pneumonia ☐ May be presented with fever, headache, myalgia and diarrhea. or bronchopneumonia >Mycoplasma pneumoniae: mostly presented with extrapulmonary manifestations SUCh as More with healthy young adults ☐ Myringitis inf. Tympanic membrane (ear drum) ☐ Encephalitis ■ Myocarditis S. aureus CAP- Dangerous Sever pneumonia with diffuse nodule ■ Not common ☐ Post Influenza complication. ☐ Compromised host, Co-morbidities, Extreme of age ☐ May be MSSA or MRSA (community acquired MRSA) ☐ Multi lobar involvement, necrosis of lung with cavitations causing lung abscess or multiple pyemic abscesses and empyema ☐ Septic Arthritis ☐ Hypoxemia, and Hypotension are common

above transverse fissur mostly (upper lope) under it (middle or lower lope)



Bilateral opacity in middle and lower lope (multilopar pneumonia



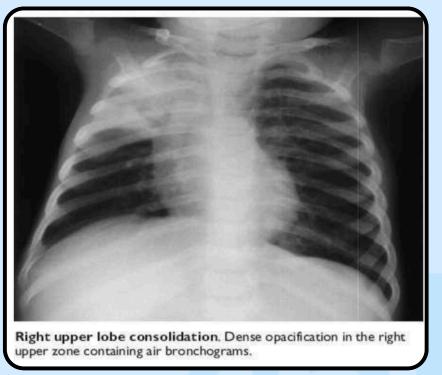
Appearnce in X-Ray help you to detect severity

Rt Upper lope pneumonia

Chest X-ray

Normal chest film Posteroanterior view of a normal chest radiograph. Courtesy of Carol M Black, MD.

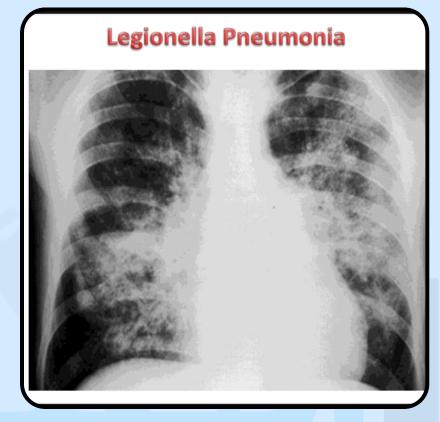
Diagnosis, prognosis, pathogens......



Rt Upper lope pneumonia



Patchy + diffused in upper + lower zone "heterogeneous "(bronchopneumonia)

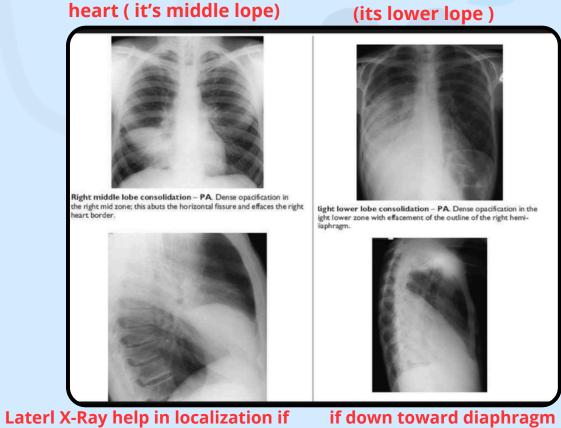


Opacity under fissure joined with Rt border of heart (it's middle lope)

opacity extreme above toward

if lean to heart (middle lope)

sternum (it's upper lope)

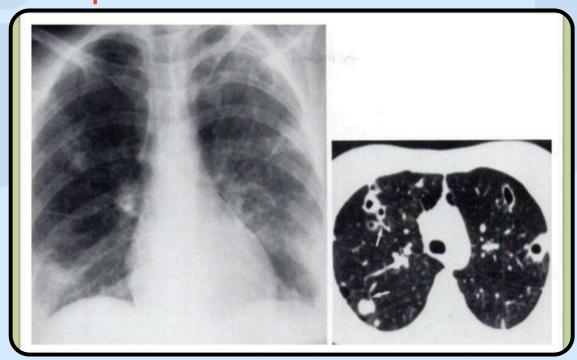


(lower lope)

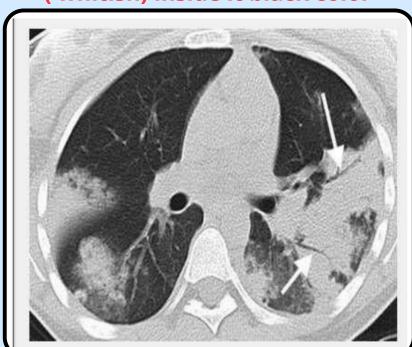
Opacity under fissure

joined w/ Diaphragm

In staph.A diffused nodules with cavitation in CT

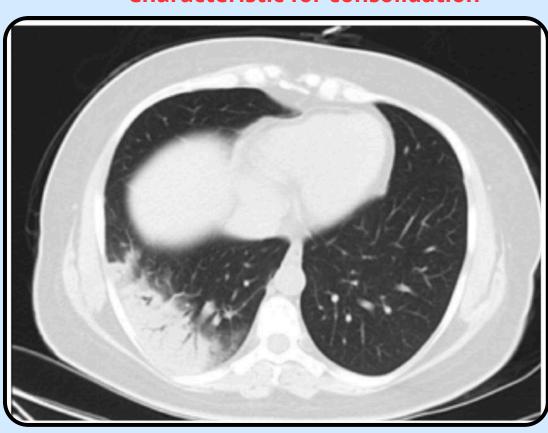


Consolidation ear bronchogram (whitish) inside it black color



Same

Characteristic for consolidation



When patient need hospitalization

Laboratory Tests for CAP

• CBC.

- ABG
- Gram stain of sputum.
- Culture of sputum. • Blood cultures.
- Liver enzymes
- CRP and ESR
- BUN and Creatinine
- Serum electrolytes
 - زي الخيوط

Complications of CAP

- Parapneumonic effusion. Accumulating fluid
- Empyema
- in plural cavity
- Lung abscess- destruction of lung .
- Multiple Pyaemic Abscesses
- Septicemia Brain abscess, Liver Abscess
- Hypotension and septic shock

Risk Factors for Hospitalization in CAP

- Old Age
- Comorbidities:
- Chronic chest diseases
- ➤ Asthma,
- >COPD,
- > Bronchiectasis
- Chronic diseases:
- ➤ Diabetes,
- >CHF,
- **≻**Neoplasia

Criteria for severe pneumonia:

Minor criteria

If patient has 3=/< sever pneumonia, admit to ICU

- 1. Confusion
- 2. Respiratory rate> 30 breaths/min
- 3. Hypothermia (core temperature, <36C)
- 4. Hypotension requiring aggressive fluid resuscitation
- 5. Multilobar infiltrates
- 6. Leucopenia: (WBC <4000 cells/mm3)
- 7. Thrombocytopenia (<100,000 cells/mm3)
- 8. Uremia (BUN level, 20 mg/dL)
- 9. PaO2/FiO2 ratio < 250

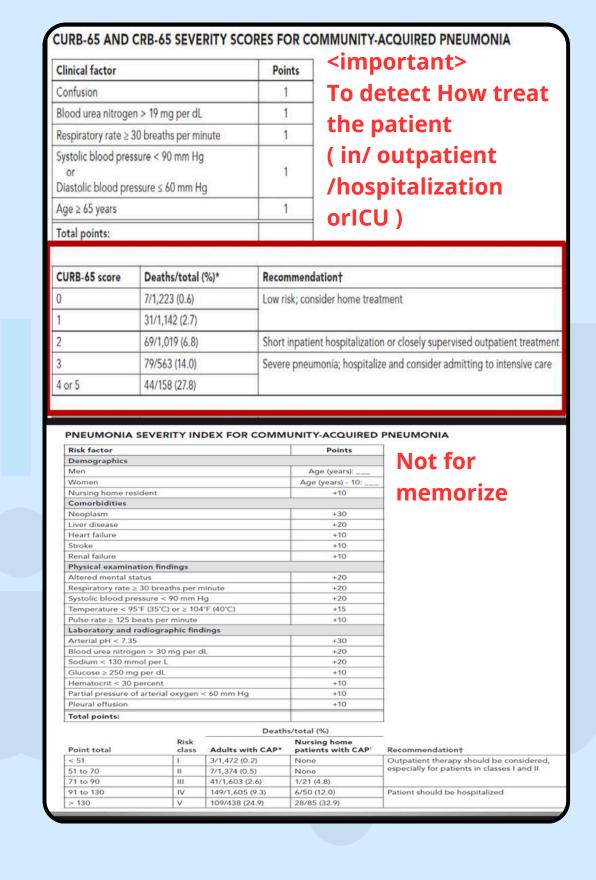
Patient Arterial O2 from ABG devide on intake (ventilated) O2

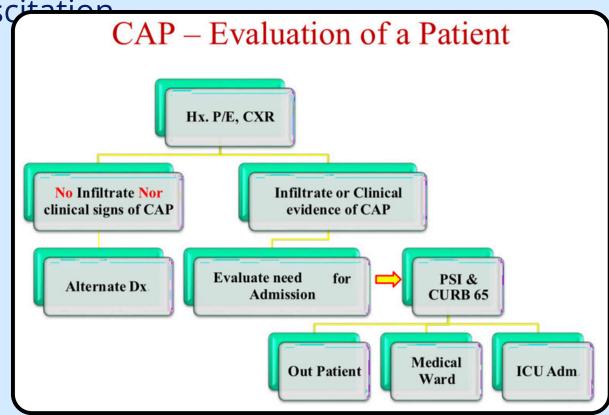
Major criteria

If has one just admit to ICU

- 1. Invasive mechanical ventilation
- 2. Septic shock with the need for vasopressors
- ICU admission = one major or 3 minor

Infiltrate Patterns and Pathogens	
CXR Pattern	Possible Pathogens
Lobar	Strept. Pneumoniae.Gram neg. e.g. Klebsiella, H. influ.
Patchy	Atypical, Viral
Interstitial	Viral, Legionella
Cavitatory	S.aureus, Klebsiella, AnaerobesTB, Fungi
pleural effusion	StaphKlebsiella





Treatment

Patients should initially be treated empirically, based on the likely pathogens for each patient group.

Supportive measures

- a. Bed rest.
- b. Adequate nutrition either orally or IV in severe cases.
- c. Fluid and electrolytes replacement.
- d. Analgesics for pain and antipyretics for fever.
- e. Respiratory support by oxygen supply or mechanical ventilation.
- f. Circulatory support by inotropic agent in hypotension.
- g. Steroids may be used to suppress the inflammatory response to infection. CAP

The patient may be Previously Healthy or has Comorbidities such as:

- Chronic heart, lung, liver, and renal disease,
- Diabetes mellitus, Alcoholism,
- Malignancies, Immunosuppression
- Asplenia,
- Use of antimicrobials within the previous 3 months

Group I: Outpatients but no Comorbidities Organisms One of these Therapy Streptococcus pneumoniae Macrolide: - Azithromycin500mg once or Hemophilus influenza - Clarithromycin 500mg bid Mycoplasma pneumoniae - Erythromycin Chlamydia pneumoniae Legionella spp **or** Betalactam Amoxicillin or Respiratory viruses amoxicillin + clavulanic acid Erythromycin is not active against H. Influenza and the advanced generation Macrolides Azithromycin and Clarithromycin are better Many isolates of S. pneumoniae are resistant to tetracycline, and it should be used only if the patient is allergic to or intolerant of macrolides. **Group II: Outpatient, with Comorbidities ORGANISMS Therapy** Strept. pneumoniae (including β-Lactam; Amoxicillin, Cefpodoxime, Hemophilus influenza Cefuroxime Mycoplasma pneumoniae Amoxicillin /clavulanate Chlamydia pneumoniae Ampicillin-sulbactam Legionella spp., Ceftriaxone + Macrolide Enteric gram-negatives Aspiration(anaerobes) One of these Lung Fluroquinolones as Monotherapy Respiratory viruses

Levofloxacin 750 mg OD Moxifloxacin 400 mg OD Gemifloxacin 320 mg OD

Group III: Inpatient (Not in ICU) **ORGANISMS Therapy** Strept. pneumoniae (including DRSP) β-Lactam; Cefpodoxime, Hemophilus influenza Cefuroxime, 1.5 g bid or tds Amoxicillin /clavulanate, 1.2 Mycoplasma pneumoniae g bid or tds Chlamydia pneumoniae Legionella spp., Ampicillin-sulbactam Ceftriaxone, 1-2 gm od Enteric gram-negatives Cefotaxime Aspiration(anaerobes) Respiratory viruses Macrolide **Or** combination **B-lactam + Lung Floroquinolones Group IV: ICU- Admitted Patients** A. No Risks for Pseudomonas aeruginosa or MRSA **Therapy Organisms** Streptococcus pneumoniae Intravenous β-lactam;

Amoxicillin /clavulanate

Ampicillin-sulbactam

Intravenous Macrolide

Fluroquinolones

IV B-lactam + Intravenous

Cefotaxime

Ceftriaxone

(including DRSP)

Legionella spp.

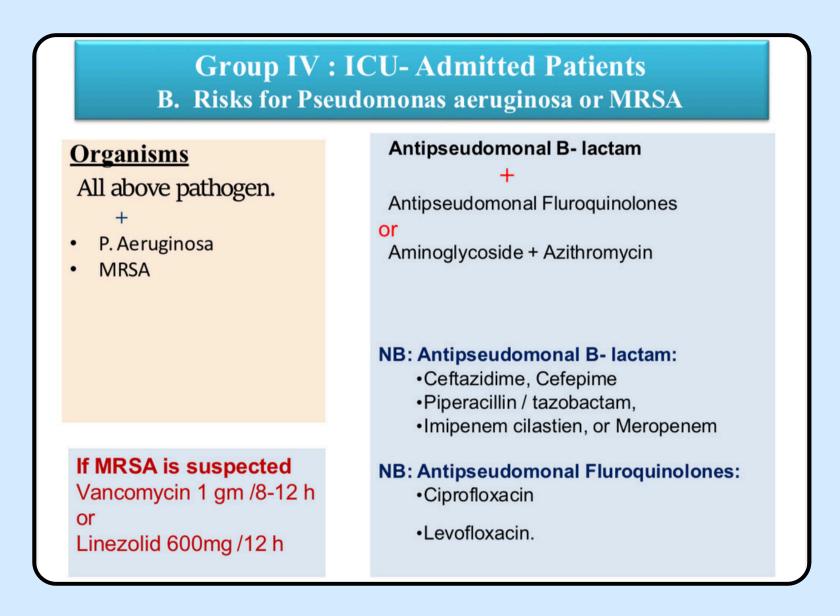
Hemophilus influenzae

Mycoplasma pneumoniae

Enteric gram-negatives

Aspiration(anaerobes)

Respiratory viruses



Anti-influenza treatment, such as oseltamivir, be prescribed for adults with CAP who test positive for Influenza in the inpatient setting, independent of duration of illness before diagnosis (strong recommendation).

Treatment with oseltamivir is associated with reduced risk of death in patients hospitalized for CAP who test positive for influenza virus

Treatment within 2 days of symptom onset or hospitalization may result in the best outcomes, although there may be benefits up to 4 or 5 days after symptoms begin Antiviral drugs against COVID-19 e.g. Favipiravir should be used in case of confirmed

Patient follow up

Patients should be evaluated after 2-3 days for initial improvement in:

- Clinical parameters e.g. Fever and toxic symptoms.
- Lab parameters e.g. leukocytosis and acute phase reactant.
- Chest radiograph findings shows no progression but usually clear within 1-4 weeks but may persist for longer duration in older individuals and those with underlying pulmonary disease

Switch to Oral Therapy

Four criteria

- 1. A febrile on two occasions 8 h apart
- 2. Improvement in cough, dyspnea & clinical signs
- 3. WBC decreasing towards normal
- 4. Functioning GI tract with adequate oral intake

Duration of Therapy

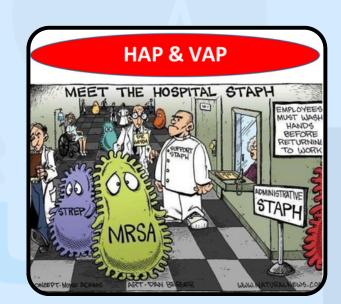
- For patients with low or moderate severity and uncomplicated pneumonia, 5-7 days of appropriate antibiotics is recommended.
- For those with high severity and complicated pneumonia or microbiologically-undefined pneumonia, 7–10 days treatment is proposed.
- If S. aureus or Gram-negative enteric bacilli pneumonia is suspected or confirmed 14 21 days treatment is used

Risk factors for treatment failure:

- 1. Age > 65
- 2. Patient with comorbidities:
- ➤ Neoplasia .
- > Liver disease.
- ➤ Neurologic disease.
- > Structural lung disease e.g. Bronchiectasis.
- 3. Multilobar pneumonia.
- 4. Cavitation, pleural effusion.
- 5. Leukopenia.
- 6. Aspiration pneumonia.
- 7. Infection with MRSA, Legionella, or gram-negative bacilli

CAP- Management summery......

- CURB-65 scoring and Classification of cases
- Sputum and Blood culture collection in the first 24 h prior to Antibiotic administration.
- Early Empirical Antibiotic administration within 4-6 hours
- Empirical on non Empiric Bases
- Change Antibiotic according to pathogen & sensitivity pattern
- Pneumococcal & Influenza vaccination; Smoking cessation specially old age or COPD



Late Infection (≥5 days) OR

Hospital acquired pneumonia (HAP)

· Defined as pneumonia that occurs 48 hours or more after admissio

Risk Factor For MDR pathogens

Antimicrobial therapy in the preceding 3 months Present hospitalization of ≥5 days

High frequency of antibiotic resistance in the community or in the specific hospital unit Hospitalization for ≥48 h in the preceding 3 months

Home infusion therapy including antibiotics

Home wound care

Chronic dialysis within 1 month
Family member with MDR pathogen
Immunosuppressive drug and/or therapy

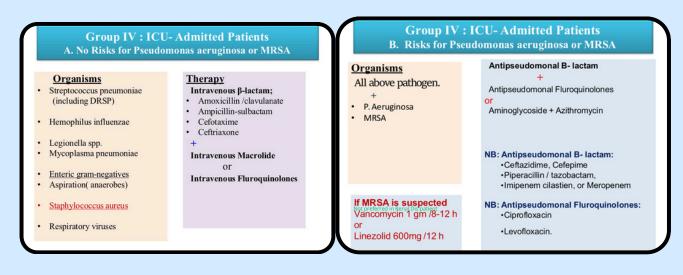


Early infection (<5 days) AND

Suspected hospital-acquired or ventilator-associated pneumonia

Send sputum/endotracheal tube aspirate and blood cultures

Deal with him as sever pneumonia



Duration of Therapy

- Optimal duration of antimicrobial therapy in HAP patients is 10-14 days.
- A trend to greater rates of relapse for short duration therapy was seen if the etiologic agent was P. aeruginosa or an Acinetobacter species, so treatment duration is 14-21 day