

Treatment of Respiratory Tract Infections

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Objectives of the lecture

- ▶ At the end of lecture , the students should be able to understand the following:
- ▶ The types of respiratory tract infections
- ▶ The antibiotics that are commonly used to treat respiratory tract infections and their side effects.
- ▶ Understand the mechanism of action, pharmacokinetics of individual drugs.

Upper respiratory tract

Nasal cavity

Pharynx

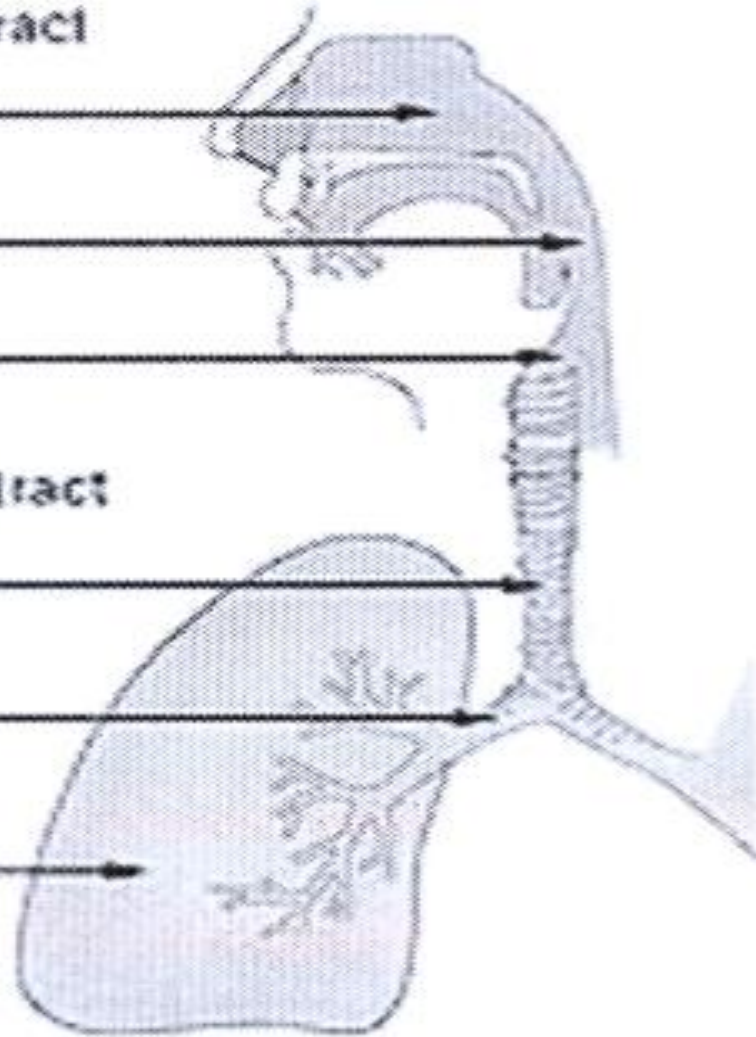
Larynx

Lower respiratory tract

Trachea

Primary bronchi

Lungs



Causes of URTI's

- ▶ **Viruses** (rhinoviruses, influenza viruses, corona viruses, etc.)

(Should not be treated with antibiotics)

Treatment: rest and plenty of fluids, OTC cold, pain relievers.

- ▶ **Bacteria** (mainly Group A streptococcus, H. influenza)

Treatment: Antibiotics. The type depends on:

Type of bacteria

Sensitivity test

LRTI's (costly & more difficult to treat)

▶ **Bronchitis** (inflammation of major trachea & bronchi)

Acute

Chronic

Acute exacerbation of chronic bronchitis

Causes: viruses or bacteria (H. influenza, S. pneumoniae & M. catarrhalis).

▶ **Pneumonia** (Serious infection of bronchioles & alveoli)

Community-acquired (CAP)

Hospital-acquired (nosocomial)

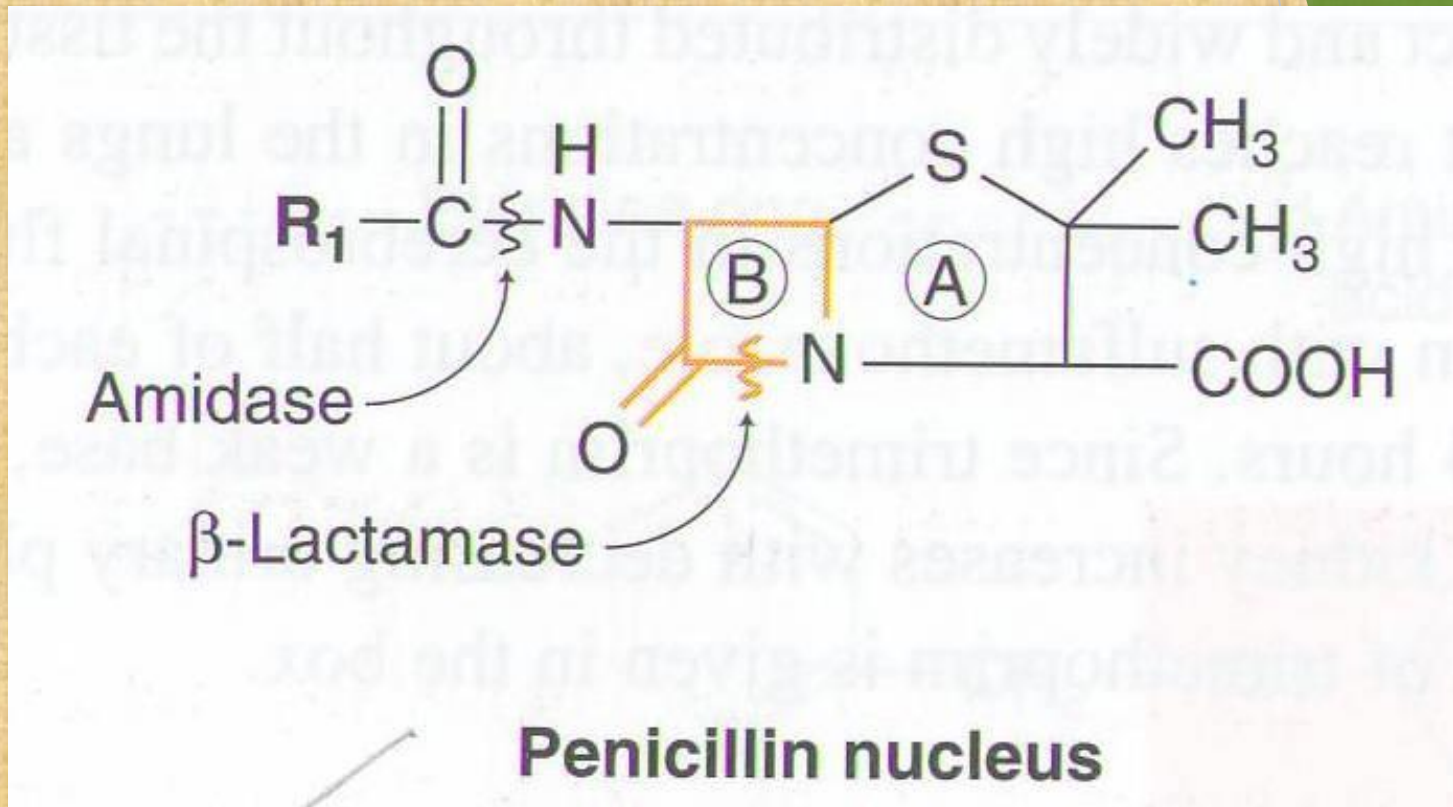
Causes: Bacteria

S. pneumoniae** (66%), H. influenza (20%), M. catarrhalis (20%)

Antibiotics commonly used in the treatment of RTI's

- ❑ **Beta lactam antibiotics
(Penicillins / Cephalosporins)**
- ❑ **Macrolides**
- ❑ **Fluoroquinolones**
- ❑ **Tetracyclines**
- ❑ **Aminoglycosides**

Penicillins



Mechanism of action of Penicillins

- ▶ Inhibits bacterial cell wall synthesis through inhibition of peptidoglycan layer of the cell wall.
- ▶ Bactericidal

Pharmacokinetics of Penicillins

- ❖ Given orally or parenterally
- ❖ Not metabolized in human.
- ❖ Relatively lipid insoluble.
- ❖ Excreted mostly unchanged in urine (glomerular & tubular secretion).
- ❖ Probenecid slows their elimination and prolong their half life.
- ❖ Half-life 30-60 min (increased in renal failure).

Broad- spectrum penicillins

- ▶ **Amoxicillin- Clavulanic acid**
- ▶ **Ampicillin- Sulbactam**
- ▶ **Piperacillin- tazobactam**

**Act on both gram+ve & gram-ve
microorganisms**

Hypersensitivity
reactions

*Adverse
effects*

Convulsions
(after high
i.v. dose or
in renal
failure)

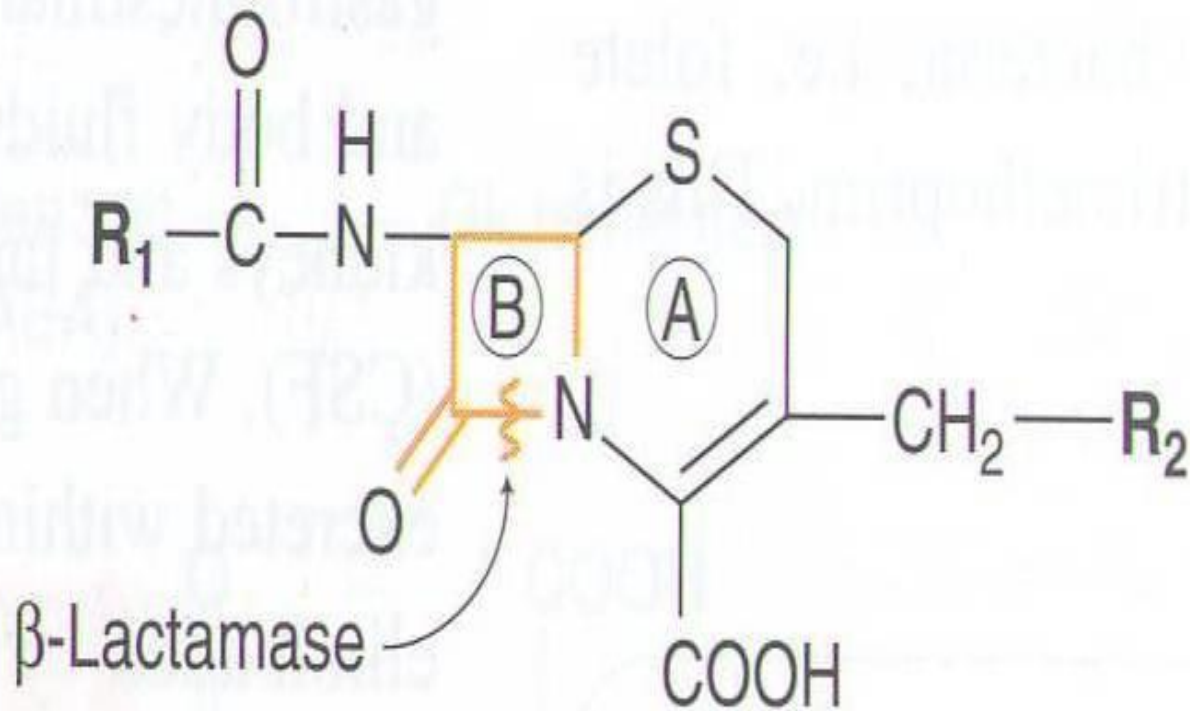
Diarrhea
Superinfections

Nephritis

Therapeutic uses of Penicillins

- ▶ Upper respiratory tract infections
- ▶ Lower respiratory tract infections

Cephalosporins



Cephalosporin nucleus

Mechanism of action of Cephalosporins

- ▶ Inhibit bacterial cell wall synthesis
- ▶ Bactericidal

(Similar to Penicillins)

Pharmacokinetics of Cephalosporins

Given parenterally and orally.

Relatively lipid insoluble (like penicillins)

Mostly excreted unchanged by the kidney (glomerular & tubular secretion).

Probenecid slows their elimination and prolong their half live.

Half-life 30-90 min; **except, ceftriaxone 4-7 hr**

1st Generation Cephalosporins

▶ Cephalexin

- Given orally
- Effective against gram positive bacteria.
- Effective in URTI's

2nd Generation Cephalosporins

Cefuroxime axetil , cefaclor

- ▶ Given orally
- ▶ Effective mainly against Gram-negative bacteria.
- ▶ Well absorbed orally
- ▶ Active against β -lactamase -producing bacteria

Uses:

- ▶ Upper and lower respiratory tract infections

3rd Generation Cephalosporins

Ceftriaxone / Cefotaxime / Cefixime

- ▶ Given by intravenous route
- ▶ More effective against gram-negative bacilli
- ▶ Effective treatment in pneumonia

Adverse effects of cephalosporins

1

- Hypersensitivity reactions

2

- Thrombophlebitis

3

- Superinfections

4

- Diarrhea

Macrolides

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graph TD; A[Macrolides] --> B[Erythromycin]; B --> C[Azithromycin]; B --> D[Clarithromycin];
```

Erythromycin

Azithromycin

Clarithromycin

Mechanism of action

Inhibit protein synthesis by binding to 50S subunit of the bacterial ribosomes

Bacteriostatic

Bactericidal at high concentrations

Clarithromycin

- ▶ **More effective on G+ve bacteria.**
- ▶ **Stable at gastric acidity**
- ▶ **Inhibits cytochrome P450 system**
- ▶ **Metabolized to active metabolite**
- ▶ **Biliary route is the major route of elimination**
- ▶ **Only 10-15% excreted unchanged in the urine**
- ▶ **Half-life 6-8 hours**

Azithromycin

- ▶ More effective on G-ve bacteria.
- ▶ Stable at gastric acidity
- ▶ Undergo some hepatic metabolism (inactive metabolite)
- ▶ Biliary route is the major route of elimination
- ▶ Only 10-15% excreted unchanged in the urine
- ▶ Half- life (3 days)
- ▶ Once daily dosing

No effect on cytochrome P- 450

Clinical uses of Macrolides

- ▶ Chlamydial pneumonia
- ▶ Legionella pneumonia

Adverse effects

GI disturbances(nausea, vomiting, abdominal cramps & diarrhea.

Hypersensitivity reactions

Fluoroquinolones

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graph TD; A[Fluoroquinolones] --- B[Ciprofloxacin]; A --- C[Moxifloxacin]; A --- D[Gatifloxacin];
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Ciprofloxacin

Moxifloxacin

Gatifloxacin

Mechanism of action

Inhibit DNA Gyrase enzyme

(an enzyme involved in DNA supercoiling)

Antibacterial spectrum

Ciprofloxacin mainly effective against G - bacteria

Moxifloxacin & Gatifloxacin G - & G + & given once daily.

(highly active against Pseudomonas species)

Pharmacokinetics

- **Given orally or parenterally.**
- **Di & tri-valent cations interfere with its absorption**
- **Concentrates in many tissues (kidney, prostate, lung & bones/ joints)**
- **Excreted mainly through the kidney**
- **Their relatively long Half-life allow once daily (moxifloxacin & Gatifloxacin) & twice-daily (ciprofloxacin) dosing.**

Clinical Uses



Acute exacerbation of chronic obstructive pulmonary disease



Community acquired pneumonia



Legionella pneumonia

Adverse effects

- ❖ Nausea , vomiting , diarrhea
- ❖ CNS effects (confusion, insomnia, headache, anxiety).
- ❖ Damage of growing cartilage(**arthropathy**)
- ❖ Phototoxicity(avoid excessive sunlight)

Contraindications

- ▶ Not recommended for patients younger than 18 years
- ▶ Pregnancy
- ▶ Breast feeding women

Tetracyclines (e.g. Doxycycline)

It is a long acting tetracycline

Mechanism of action

Inhibit protein synthesis by binding reversibly to 30 s subunit

Doxycycline (Cont.)

Pharmacokinetics

Usually given orally

Absorption is 90-100%

Absorbed in the upper s. intestine & best in absence of food

Food & di & tri-valent cations (Ca, Mg, Fe, AL) impair absorption

Protein binding 40-80 %

Distributed well, including CSF

Cross placenta and excreted in milk

Largely metabolized in the liver

Doxycycline (Cont.)

Side effects

1. nausea, vomiting ,diarrhea & epigastric pain(give with food)
2. Thrombophlebitis - i.v
3. Hepatic toxicity (prolonged therapy with high dose)
4. Brown discolouration of teeth - children
5. Deformity or growth inhibition of bones - children
6. Phototoxicity
7. Vertigo
8. Superinfections.

Contraindications of doxycycline

- ▶ Pregnancy
- ▶ Breast feeding
- ▶ Children(below 10 yrs)

Uses of Doxycycline

- ▶ Treatment of URTIs caused by *S.pyogenes*, *S.pneumonia* & *H. influenza*.

Aminoglycosides

e.g. GENTAMICIN, i.m, i.v.

- Bactericidal antibiotics
- Inhibits protein synthesis by binding to 30S ribosomal subunits.
- Poorly absorbed orally (highly charged).
- Only active against gram negative aerobic organisms.
- cross placenta.
- Excreted unchanged in urine

Gentamicin(CONT)

Adverse effects :

- Ototoxicity
- Nephrotoxicity
- Neuromuscular blocking effect

Therapeutic uses of Gentamicin

- Severe infections caused by gram negative organisms.

THANK YOU

