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OBJECTIVES

- * At the end of lecture, the students should:
- * Discuss the etiology of tuberculosis
- Discuss the common route for transmission of the disease
- * Discusses the out line for treatment of tuberculosis
- Discuss the drugs used in the first & second line

OBJECTIVES (continue)

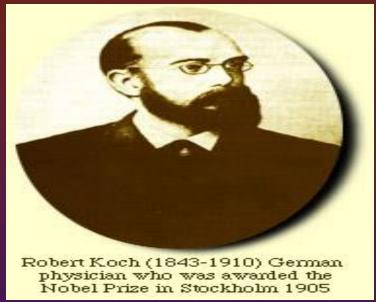
Regarding:

- ❖ The mechanism of action
- Adverse effects
- Drug interactions
- Contraindication
- Discuss tuberculosis & pregnancy
- Discuss tuberculosis & breast feeding

Etiology

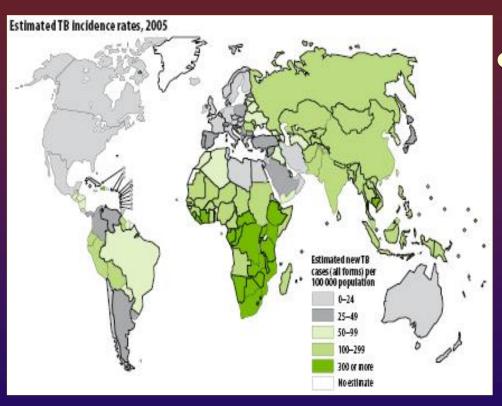
Mycobacterium tuberculosis, slow growing, an acid fast bacillus

❖ Robert Koch was the first to see Mycobacterium tuberculosis with his staining technique in 1882.



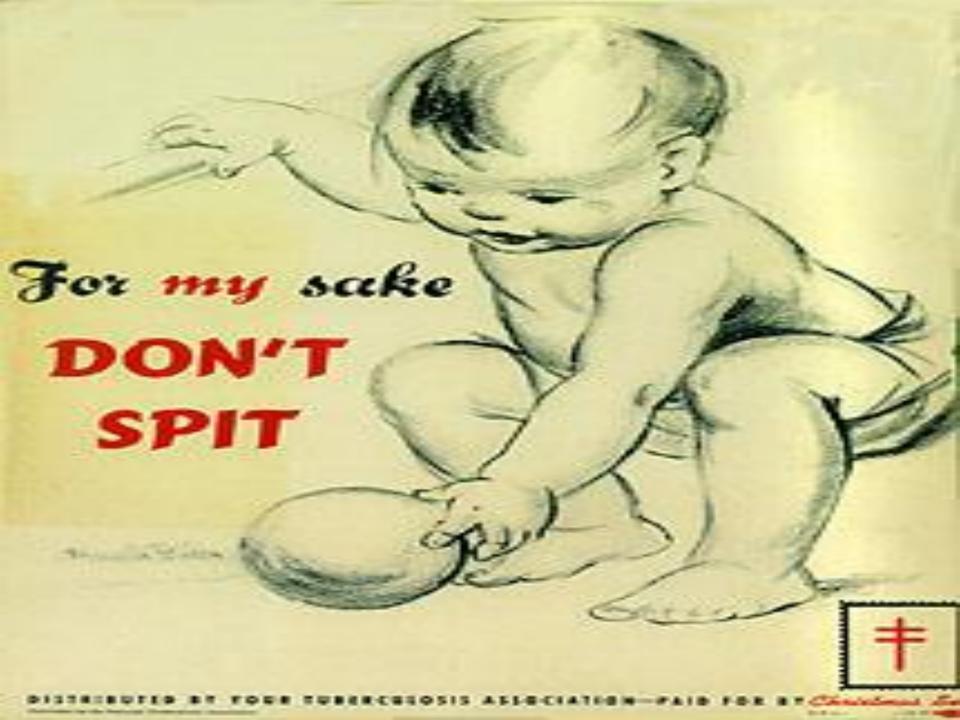


Disease information:



Each year, 1% of the global population is infected.

More than one third of the world's population has **tuberculosis**.



YOUR COUGHS AND SNEEZES



SPRAY SPREADS COLDS · FLU · TUBERCULOSIS

THE RESIDENCE THERE OF Christman Saule made the Poster Posterial

Tuberculosis

Common sites of infections

- * Apical areas of lung
- * Renal parenchyma
- ***** Growing ends of bones

Treatment Of Tuberculosis

- Preventing development of drug resistance is the most important reason to use drug combination.
- *Periods of treatment (minimum 6 months)
- Drugs are divided into two groups:
- 1. First line 2. Second line

Antimycobacterial drugs

First line

- *Isoniazid (INH)
- * Rifampin
- *Ethambutol
- * Pyrazinamide

Given for first 8 weeks, followed by INH/RIF for 18 weeks

Streptomycin (should not be the first line choice)

Never use a single drug therapy

- *Isoniazid –rifampin combination administered for 9 months will cure 95-98% of cases.
- *Addition of pyrazinamide/ethambutol for this combination for the first 2 months allows total duration to be reduced to 6 months.

Isoniazid

- *Bacteriostatic for resting bacilli.
 - *Bactericidal for rapidly growing bacilli.
 - *Is effective against intracellular & extracellular bacilli

Mechanism Of Action

Inhibits the synthesis of mycobacterial

cell wall (inhibit the synthesis of mycolic acid)

Clinical uses

- * Treatment of TB
- Treatment of Latent TB in patients with positive tuberculin skin test
- Prophylaxis against active TB in individuals who are in great risk.

Adverse effects

- *Peripheral neuritis
 - (pin & needles sensation in the feet
- *Optic neuritis & atrophy.
 - (Pyridoxine should be given in both cases)
- *Hepatitis (toxic metabolites)
- * Hepatitis with INH, is age dependent; it is rare in persons younger than 20 years, risk increases with age and alcohol use

Drug Interactions of INH

- Enzyme inhibitor
- *Slow and fast acetylators.



- *Bactericidal
- *Inhibits RNA synthesis
 by binding to DNA dependent RNA

polymerase enzyme.

Site of Action (similar to INH)

- *Intracellular bacilli
- *Extracellular bacilli

Clinical uses

*Treatment of TB

*Prophylaxis.

Adverse effects

- *Harmless red-orange discoloration of body secretions (saliva, sweat, tears). Tell the patient about this effect. Can permanently stain contact lenses.
- *Hepatitis less common compared to INH
- Flu-like syndrome
- *Hemolytic anemia

Drug Interactions

- Enzyme inducer
- Clincally significant drug interactios such as warfarin, methadone will be metabolized faster

Ethambutol

*Bacteriostatic

*Inhibitor of mycobacterial arabinosyl transferase (alters the cell barrier) disrupts the assembly of mycobacterial cell wall.

Site Of Action (similar to INH)

*Intracellular & Extracellular bacilli

Clinical uses

*Treatment of tuberculosis in combination with other drugs.

Adverse effects

*Impaired visual acuity

*red-green color blindness.

 Ethambutol is contraindicated in children under 5 years.

Pyrazinamide

- *Bacteriostatic
- *Mechanism of action is unknown.

Site Of Action

* Active against Intracellular Bacilli

Clinical uses

- *Mycobacterial infections mainly in multidrug resistance cases.
- *It is important in short –course (6 months) regimen.
- *Prophylaxis of TB.

Adverse effects

*Hepatotoxicity (common)

*Hyperuricemia (gouty arthritis)

Drug fever & skin rash

Streptomycin

- *Bactericidal
- *Inhibitors of protein synthesis by binding to 30 S ribosomal subunits.
- *Active mainly on extracellular bacilli

Clinical uses

*Severe, life-threating form of T.B. as meningitis, disseminated disease.



- Ototoxicity
- *Nephrotoxicity
- *Neuromuscular block

Indication of 2nd line treatment

- *Resistance to the drugs of 1st line.
- *Failure of clinical response
- *There is contraindication for first line drugs.
- * Used in typical & atypical tuberculosis
- * 2nd line drugs are more toxic than 1st line drugs

Ethionamide

*Inhibits the synthesis of mycolic acid

Clinical uses

*As a secondary line agent ,treatment of TB.

Adverse Effects

Terratogenic

Poorly tolerated

Because of:

- *Severe gastric irritation &
- *Neurological manifestations

Fluoroquinolones (Ciprofloxacin)

***Effective against multidrug- resistant tuberculosis.**

Rifabutin

- * RNA inhibitor
- Cross –resistance with rifampin is complete.
- *Enzyme inducer

Clinical uses

*Effective in prevention &treatment of T.B.

*In prevention & treatment of atypical TB.

Adverse Effects

***GIT** intolerance

***Orange-red discoloration of body secretions.**

Aminosalicylic Acid (PAS).

*Bacteriostatic

*Inhibits Folic acid synthesis.

Clinical uses

*As a second line agent is used in the treatment of pulmonary & other forms of tuberculosis.

Adverse effects

*GIT upset

* Crystalluria

TB & Pregnancy

- Untreated TB represents a great risk to the pregnant woman & her fetus than the treatment itself.
- * First line (INH, Ethmabutol and rifampicin) drugs are given for 9 months in normal doses
- Streptomycin not used

TB & Breast Feeding

* It is not a contraindication to receive drugs, but caution is recommended

