

# ***TUBERCULOSIS***

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mankind.

- Causative organism -Mycobacterium
- Etiopathogenesis of the disease is clearly understood.
- A vaccine against Tuberculosis has been available for close a century.

## Introduction

development.

- ***An overwhelming majority of cases*** and practically all deaths due to Tuberculosis take place in developing countries.
- *Tuberculosis is currently second only to AIDS* as an infectious cause of death worldwide.

***falling globally since 2003 .***

- Organism can infect practically any organ of the body.
- Pulmonary tuberculosis accounts for over 80% of the total cases suffering from tuberculosis.

- meningeal,
- bone and joint,
- renal,
- genital,
- abdominal or mesenteric and
- tubercular lymphadenopathy .

□ Tuberculosis is the ***commonest opportunistic infection in patients suffering from AIDS*** in large parts of the world.

- Transmitted by droplet nuclei after close contact with a person who has infectious disease.
- Effective treatment - available for over sixty years.
- World Health Organization declared TB as a ***global public health emergency in 1993.***
- Treatment : prolonged multidrug therapy which *increases the potential risk of non-adherence by patients*

• TB known by a number of names through

- Pott's disease: TB of the bone with characteristic vertebral fusion and deformity of the spine.
- The most familiar term for TB was consumption, which means to consume or wear away.

# HISTORY

- '***Captain of the Men of Death***': the most fitting name among all
- Egyptian mummies from 2400 BC: show ***definite pathological signs of tubercular decay in fragments of the spinal column.***
- Around 460 B.C., Hippocrates identified phthisis as the most widespread disease of the times, and noted that it was almost always fatal.
- ***Sylvius was the first to identify actual tubercles in the lungs*** and other areas of consumptive patients in 1679.
- In 1882, Robert Koch discovered a staining technique that enabled him to see *Mycobacterium tuberculosis*.



- Over the centuries, TB has taken over 1 billion lives.
- Deaths due to tuberculosis comprise 25% of all avoidable deaths in developing countries.
- ***95% of TB cases and 98% of TB deaths*** are in developing countries.
- 75% of TB cases in developing countries are in the ***economically productive*** age group of 15 - 50 years.

## **Epidemiology: Global**

- approximately 1 billion people will be newly infected,
- over 150 million will get sick, and
- 36 million will die of TB,

***if control is not further strengthened***

- 1.8 million cases occurring annually,
- Accounts for a fifth of the world's new TB cases and 2/3<sup>rd</sup> of the cases in South - East Asia.
- This makes India the *highest TB burden country in the world*

## Epidemiology: India

- Because it affects adults, tuberculosis causes ***enormous social and economic disruption.***
- More than 80% of the burden of tuberculosis is ***due to premature death,*** as measured in terms of Disability Adjusted Life Years (DALYs) lost.

- 54 years).
- The burden of TB is enormous but is hidden by stigma.
- TB kills ***more women in India*** than any other infectious disease.
- Women with tuberculosis are often ***severely stigmatized***.

# Agent :

## **Mycobacterium tuberculosis**

- genus Mycobacterium,
- family Mycobacteriaceae and Order Actinomycetales.
- Gram positive, non - motile, non - sporing, pleomorphic rod.
- Bacilli are obligate aerobes growing most successfully in tissues having the highest partial pressure of oxygen, such as lung apices.

hours. Hence, lesions typically evolve in a sub - acute to chronic course.

- They are classified as *Acid - Fast Bacilli (AFB)* because they retain the carbol - fuchsin red dye after washing with acid, alcohol, or both.
- *Mycobacterium bovis* is the etiologic agent of TB in cows and rarely in humans.
- Both cows and humans can serve as **reservoirs**.

• Humans can also be infected by the

- *Mycobacterium africanum* can be a rare cause of tuberculosis.

Other human pathogens belonging to the genus *Mycobacterium* include

- ***Mycobacterium avium*** which causes a TB - like disease especially prevalent in AIDS patients and
- *Mycobacterium leprae*, the causative agent of leprosy.



- Nutrition
- Immunity

**Host factors**

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- Overcrowding
- Undernutrition
- Population explosion
- Lack of education
- Large families

**Social factors**

and droplet nuclei.

- Incubation periods: weeks, months or years.

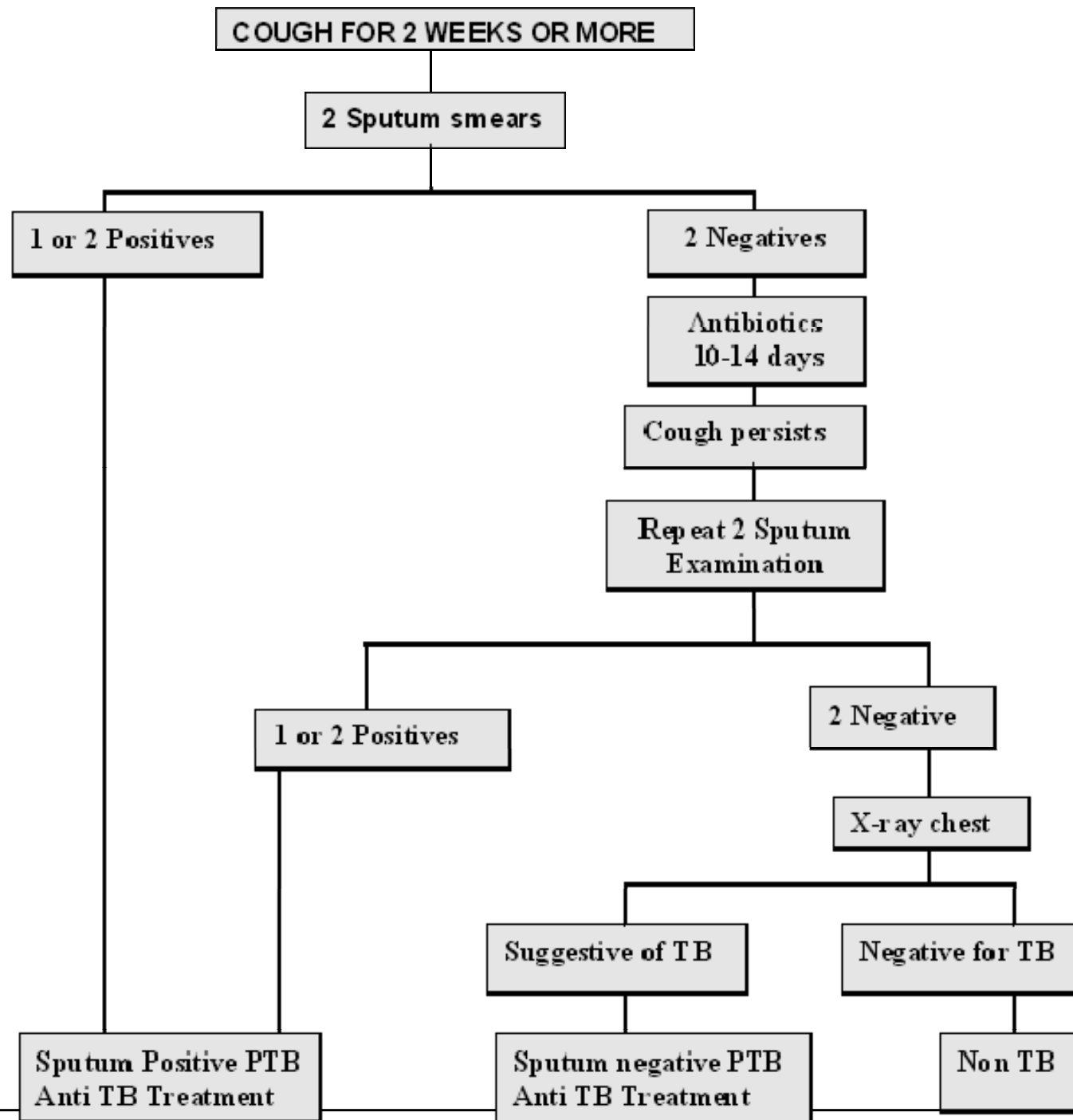
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culture

- Treatment
- BCG vaccination
- Chemoprophylaxis
- Rehabilitation
- Nutrition

## **Prevention & Control**

# Diagnostic Algorithms for Pulmonary TB



# Revised Categories

Treatment groups	Type of patient	Regimen	
		Intensive phase (IP)	Continuation phase (CP)
<b>New (Cat I)</b>	<ul style="list-style-type: none"> <li>•New sputum smear positive</li> <li>•New sputum smear negative</li> <li>•New extra-pulmonary</li> <li>•New others</li> </ul>	2 H3R3Z3 E3	4 H3R3
<b>Previously treated (Cat II)</b>	<ul style="list-style-type: none"> <li>•Smear positive relapse</li> <li>•Smear positive failure</li> <li>•Smear positive treatment after default</li> <li>•Others</li> </ul>	2 H3R3Z3 E3S3/ 1 H3R3Z3	5 H3R3E3

# MDR-TB and DOTS-Plus

- MDR-TB is a *lab diagnosis*, NOT a clinical one
- MDR-TB levels of less than 1% to 3% in new cases and of *12% in re-treatment cases*.
- Emergence of resistance to Rifampicin in only 2% of patients, despite a high level (8%) of initial resistance to Isoniazid, either alone or in combination with other anti-TB
- *Quality assured laboratory facility for culture* and *Drug Susceptibility Test* must be available (NB: 2 – 4 months delay before DST results seen)

- Category II patient who is smear positive at the ***end of the fourth month of treatment or later*** will be identified as “**MDR-TB suspect**” and will be tested by culture and DST
- A patient who is an “MDR TB Suspect” should be referred by the respective medical officer – peripheral health institute (MO-PHI) to the District TB Officer (DTO)



- An MDR-TB suspect who is sputum culture positive and whose TB is due to bacilli that are ***resistant in-vitro*** to at least isoniazid and rifampicin (the DST result being from an RNTCP accredited IRL).

- **24 months standardized ZHD HNC regimen given under **daily** DOT:**
  - **6 Km Ofx Eto Cs 7 F / 18 Ofx Eto Cs F**

- Dosages based on  $\leq 45\text{kgs}$  /  $>45\text{kgs}$  wt bands
- Drugs supply: 3 months in IP / 6 months in CP
- All patients to receive pyridoxine 100mg daily
- *PAS to be used as substitute drug in case of severe ADRs* , crucial to monitor for adverse drug reactions and treat when required.

# HIV & TB

- HIV co-infection strongest known risk factor for the progression of latent TB infection to active TB disease
- Estimated 7-10% annual risk of reactivation, with 60% lifetime risk (cf. 10% lifetime risk in TB infected, non-HIV infected individual)
- Conversely, TB amongst the most common causes of morbidity and mortality in people living with HIV/AIDS
- Immune response to TB bacilli increases HIV replication leading to a rapid progression of HIV disease
- Optimal access to DOTS will significantly reduce