



Tuberculosis (TB) is a contagious airborne disease, caused by inhalation of a bacterium called *Mycobacterium tuberculosis*, that mainly affects the lungs.

TUBERCULOSIS^{1, 2}

- Tuberculosis is recognized as a major global health problem and one of the leading causes of death linked to a single infectious agent.
- Main countries concerned are low- and middle-income countries due to poverty and lack of access to proper sanitation.
- Seven countries account for 64% of TB-related deaths: India, Indonesia, China, Philippines, Pakistan, Nigeria and South Africa.
- Multidrug-resistant TB (MDR-TB) remains a public health crisis and a health security threat. A global total of 206,030 people with multidrug-resistant TB were detected and notified in 2019, a 10% increase from 2018.
- Ending the TB epidemic by 2030 is one of the health targets of the United Nations Sustainable Development Goals (SDGs).

TRANSMISSION^{1,3}

- TB spreads through inhaling tiny droplets from the coughs or sneezes of a person with active TB disease (1 person can infect 15 others).
- Poverty and poor living conditions (overcrowding, lack of ventilation) lead to increased transmission of *Mycobacterium tuberculosis*.
- Mainly inter-human transmission (rare cases of bovine transmission).

THE BURDEN OF TUBERCULOSIS¹

1/4 of the global population is infected with *Mycobacterium tuberculosis*, presenting a latent TB form, of which 10 to 15% will progress to active disease.

10 million people develop active TB disease each year

1.4 million people die annually from TB

>95% of TB deaths occur in LMIC* countries

64% of TB-related deaths occur in 7 countries

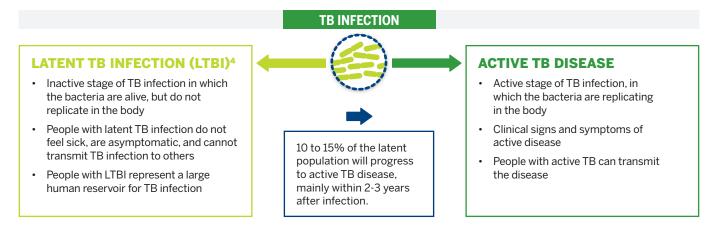


*LMIC: low- and middle-income countries

TUBERCULOSIS INFECTION¹

Tuberculosis has 2 major forms: latent TB infection (LTBI) and active TB disease.

- 90 to 95% of people infected with TB develop immunity and do not transmit infection. This form is known as latent TB infection.
- 5 to 10% of people infected will develop active TB disease.



Diagnosing people with is important to prevent progression to active TB disease and stop the spread of TB.

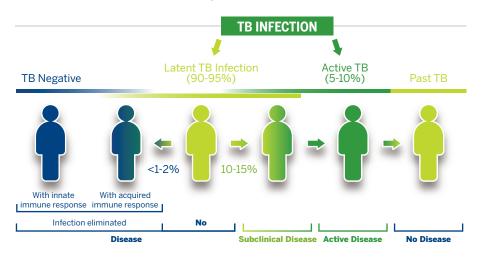


TUBERCULOSIS

STAGES OF TUBERCULOSIS INFECTION^{5, 6}

Tuberculosis infection is represented by a spectrum of stages.

Between the two main forms (latent and active), subclinical stages have been described.



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RISK GROUPS FOR LTBI^{1,3}

People at risk of being infected but with LOW RISK OF PROGRESSION to active TB disease:

- Health-care workers
- Contact of patients with active TB, IF the person is >5 vears old
- People living in communities, such as prisoners or homeless •
- Drug users •

LTBI people at HIGH RISK OF PROGRESSION to active TB disease (preventive treatment can be considered):

- Contact of patients with active TB, IF the person is <5 years old
- People living with HIV
- People receiving dialysis or organ and hematological • transplantation
- People receiving anti-TNF treatment
- People with silicosis •

Other risk factors can be associated with progression from LTBI to active TB disease: aging, poor living conditions and diabetes.⁷

CLINICAL PRESENTATION OF ACTIVE TB DISEASE*2

· Prolonged cough Fever/chills

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- Chest pain
 - Blood in sputum
- Weakness/fatigue

• Weight loss/loss of appetite

· Night sweats

*Only active TB disease is symptomatic, persons with LTBI remain asymptomatic.

DIAGNOSTIC APPROACH⁸

Diagnosis is based on:

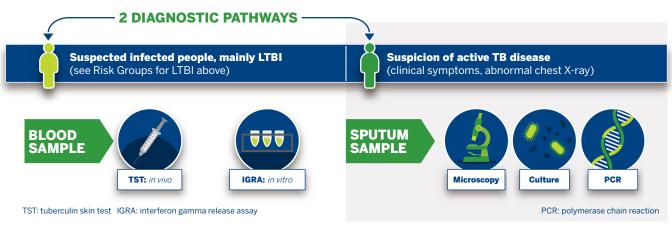
- Relevant epidemiological context (endemic region, potential exposure, proven contact with index case...)
- Anamnesis

- Clinical signs and symptoms
- Imaging: chest X-ray... .
- Laboratory testing on blood and sputum samples



TUBERCULOSIS

LABORATORY CONFIRMATION^{7,8}



Indirect diagnosis based on host response

There is NO gold standard for diagnosis of LTBI.

- **Tuberculin skin test (TST)** was the first tool used for detection of TB infection:
 - requires two doctor's visits (injection and reading 48-72 hours later)
 - reaction measurement is subjective
 - inexpensive, but lacks sensitivity and specificity (crossreaction with BCG vaccination and non-tuberculous mycobacteria (NTM))
- Recently, interferon gamma release assays (IGRA) have been developed, which measure the release of interferon gamma produced by T-cells after stimulation by specific TB antigens. IGRA are now used more often than TST, especially in high income countries:
 - require only one visit
 - objective laboratory result
 - much more sensitive and specific (no cross-reactivity with BCG and very few with NTM)
- Neither TST nor IGRA are able to distinguish between active TB and LTBI, nor predict risk of LTBI progression to active TB.
- Both assays are negatively impacted by immune depression (e.g. HIV co-infection).

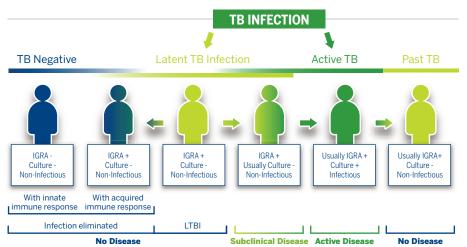
Direct diagnosis with pathogen detection/identification

- Culture from sputum specimen is the gold standard for active TB diagnosis.
- Microscopy on sputum sample remains the only diagnostic tool in many low income countries despite low sensitivity and specificity, being time-consuming and requiring skilled technicians.
- **Molecular biology** is increasingly used and WHO recommends its implementation in microscopy centers.

Antimicrobial susceptibility testing (AST)

- The gold standard for AST remains phenotypic analysis based on positive culture.
- New approaches based on genotypic assays are now emerging:
- PCR and Line Probe Assays (LPA): mixing identification of strains and prediction of resistance to major antibiotics
- Whole genome sequencing (WGS): a promising approach providing a complete picture of the bacterial identification and resistance profile

LABORATORY RESULTS ACCORDING TO TB INFECTION STAGES^{5,6}



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TUBERCULOSIS

TREATMENT¹⁰

LATENT TB INFECTION

Preventive antibiotic treatment for people at risk of progressing to active TB disease.

- Current treatment: isoniazid (9 months)
- Proposed new regimen: rifampin (4 months)

ACTIVE TB DISEASE

Active TB is never treated with a single antibiotic in order to limit the emergence of TB drug resistance.⁹ Lack of treatment compliance is also a major cause of the emergence of resistance.¹⁰

Sensitive strain

- Four drug regimen for 8 weeks: rifampin, isoniazid, ethambutol, pyrazinamide
- Followed by two drug regimen for additional 18 weeks: rifampin, isoniazid
- Resistant strain
 - Up to 2 years with second-line antibiotics: para-aminosalicyclic acid, cycloserine, ofloxacin, amikacin, etc.
- Two new drugs validated
- bedaquiline (2012), delamanid (2013)
- Two drugs under evaluation
 - linezolid and pretomanid (2019)

TB DRUG RESISTANCE9, 10

Resistance to TB antibiotics is a major obstacle to effective TB care and prevention globally.²

Multidrug-resistant TB (MDR-TB) is defined as resistance to one of the first-line antibiotics used for treatment.

Extensively drug-resistant TB (XDR-TB) is defined as resistance to first- and second-line antibiotics.

VACCINATION¹¹

Bacille Calmette-Guérin (BCG) vaccine:

- Initially designed against tuberculous meningitis (newborns & children)
- Limited protection after 10-15 years post vaccination
- Since 2006, attenuated strain of *M. bovis*: BCG SSI®

The Tuberculosis Vaccine Initiative (TBVI) is continuously working on the development of new TB vaccine candidates.

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