

Tuberculosis in the 21st Century: Challenges and Breakthroughs in Treatment

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Abstract

Tuberculosis (TB) continues to be a major global health challenge in the 21st century, with millions affected annually. While progress has been made in the treatment and management of TB, the disease remains a leading cause of death worldwide. The emergence of drug-resistant strains, delays in diagnosis, and the burden of TB in low- and middle-income countries exacerbate the global TB crisis. However, recent breakthroughs in diagnostic tools, new drug therapies, and vaccine development are offering hope for better treatment outcomes. This article explores the current challenges in TB treatment and highlights the innovative approaches and breakthroughs that could reshape the future of TB care, emphasizing the need for continued research and global cooperation in tackling this persistent threat.

Keywords: Tuberculosis, Treatment, Drug resistance, Diagnostic breakthroughs, Vaccine development, Global health, Public health, MDR-TB, XDR-TB, TB control

Introduction

Tuberculosis (TB) has been a major global health threat for centuries, causing widespread illness and death. Despite the advent of antibiotics and advances in medical science, TB remains a leading infectious cause of death worldwide, with an estimated 10 million people falling ill with TB and 1.5 million dying from it each year, according to the World Health Organization (WHO). While TB is a treatable disease, a range of challenges continue to hinder progress in its control, including the rise of multidrug-resistant (MDR-TB) and extensively drug-resistant tuberculosis (XDR-TB), delays in diagnosis, and socio-economic barriers to access to care.

The 21st century has brought some hope with the development of new diagnostic technologies, novel treatment regimens, and ongoing research into TB vaccines. This article will explore the current challenges in TB treatment, as well as the breakthroughs that are being made in diagnosis, drug therapy, and prevention, highlighting how these advances can shape the future of TB control and treatment [1].

Discussion

Challenges in Tuberculosis Treatment

Drug Resistance: A Growing Threat: The emergence of drug-resistant strains of TB has become one of the most significant challenges in the fight against the disease. Multidrug-resistant TB (MDR-TB) occurs when the bacteria become resistant to at least the two most powerful anti-TB drugs, isoniazid and rifampicin. Extensively drug-resistant TB (XDR-TB) is even more dangerous, as it is resistant to a broader range of drugs, including fluoroquinolones and injectable second-line medications. The development of MDR-TB and XDR-TB is primarily due to inadequate or incomplete treatment regimens, which result in the survival of drug-resistant bacteria [2]. In many parts of the world, particularly in low- and middle-income countries, treatment for these drug-resistant strains is often unavailable or unaffordable, leading to treatment failure and the continued spread of resistance.

Treating MDR-TB and XDR-TB is complex, costly, and often results in poor patient outcomes. Current regimens for these resistant strains can last up to two years, require the use of second-line drugs with significant side effects, and have much lower success rates than

conventional TB treatments. This poses a serious challenge to global efforts to eliminate TB [3].

Late Diagnosis and Delayed Treatment: TB is often a silent disease, with symptoms that may not appear until the disease has progressed significantly. This delayed onset of symptoms means that individuals may unknowingly spread the disease to others long before diagnosis. Additionally, in resource-poor settings, access to diagnostic tools remains limited, and diagnostic processes can be slow and cumbersome [4].

While the traditional method of diagnosing TB involves sputum microscopy and chest X-rays, these tests are not always accurate or accessible in rural and under-resourced areas. False negatives are common, leading to misdiagnosis or delayed diagnosis. Even with newer diagnostic methods, such as the GeneXpert test, which provides faster and more accurate results, the capacity to implement such technologies universally remains a challenge. Late diagnosis leads to delayed treatment, which allows the disease to spread further within communities, making TB harder to control. Early and accurate diagnosis is critical to prevent the transmission of the disease and improve treatment outcomes [5].

Socio-economic and Healthcare System Barriers: TB disproportionately affects the world's poorest populations, where overcrowded living conditions, poor nutrition, and inadequate access to healthcare exacerbate the spread of the disease. In many parts of the world, the healthcare infrastructure is insufficient to handle the burden of TB, with limited access to diagnostics, treatment, and follow-up care.

Socio-economic barriers also contribute to the persistence of TB. Many individuals with TB may face stigma or discrimination,

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particularly in communities where TB is associated with poverty or malnutrition. This stigma can discourage people from seeking treatment, further perpetuating the spread of the disease. Additionally, poverty and lack of healthcare access can prevent individuals from completing long and difficult TB treatment regimens, leading to treatment failure and the development of drug-resistant strains [6].

Breakthroughs in Tuberculosis Treatment and Diagnosis

Advances in Diagnostic Technology: Significant progress has been made in the development of rapid and accurate diagnostic tools for TB. The introduction of the GeneXpert MTB/RIF test, which detects TB and rifampicin resistance in a matter of hours, has revolutionized diagnosis in resource-limited settings. This molecular diagnostic tool has helped detect TB in individuals with limited access to traditional testing, enabling earlier treatment and reducing transmission. Additionally, advances in imaging technologies, such as digital X-rays and artificial intelligence (AI)-assisted radiology, are being explored to improve the accuracy of TB diagnosis and the detection of lung abnormalities. These technologies could reduce diagnostic delays and provide more efficient screening, particularly in high-burden settings [7].

New Drug Therapies: The treatment of TB, particularly MDR-TB and XDR-TB, has been a longstanding challenge due to the limited availability of effective drugs. However, new medications are being developed to address drug resistance and improve treatment outcomes. Bedaquiline and delamanid, two new drugs approved by the WHO for the treatment of MDR-TB, have shown promising results in clinical trials and offer hope for more effective regimens. Furthermore, new drug combinations and shorter treatment regimens are being tested to simplify and expedite the treatment process. For example, the 9-month regimen involving bedaquiline, pretomanid, and linezolid has demonstrated high success rates in the treatment of drug-resistant TB, significantly reducing treatment duration and side effects [8].

TB Vaccine Development: The Bacillus Calmette-Guérin (BCG) vaccine has been in use for nearly a century, but its effectiveness against adult pulmonary TB remains limited. The development of new vaccines has therefore become a major focus of TB research. Several promising candidates are in clinical trials, including the M72/AS01E vaccine, which has shown promising results in preventing TB infection in high-risk populations. A successful vaccine could be a game-changer in the fight against TB, particularly in preventing new infections and reducing transmission rates. Researchers are also exploring strategies to boost the immune response in individuals who have been previously vaccinated with BCG to enhance its efficacy [9].

Global Cooperation and Future Directions

The fight against tuberculosis requires global cooperation, sustained funding, and a multifaceted approach. International organizations, governments, and NGOs must work together to improve access to

diagnostics, treatment, and prevention measures, particularly in high-burden countries. Increased funding for TB research is crucial to accelerate the development of new drugs, vaccines, and diagnostic tools. Furthermore, addressing the social determinants of health such as poverty, poor nutrition, and overcrowded living conditions—will be essential to breaking the cycle of TB transmission. Improving healthcare infrastructure and reducing stigma associated with TB will also play a key role in reducing the disease burden [10].

Conclusion

Tuberculosis remains a global health challenge in the 21st century, with drug resistance, delayed diagnosis, and socio-economic factors continuing to hinder progress. However, recent breakthroughs in diagnostic technology, drug therapy, and vaccine development offer hope for improved treatment outcomes and better control of the disease. To effectively combat TB, a comprehensive approach that combines medical innovation, improved access to healthcare, and global cooperation is essential. By addressing both the medical and socio-economic factors contributing to the persistence of TB, the world can make significant strides toward eliminating this deadly disease and improving global public health.

References

1. Yagupsky P, Peled N, Riesenber K, Banai M (2000) Exposure of hospital personnel to *Brucella melitensis* and occurrence of laboratory-acquired disease in an endemic area. *Scand J Infect Dis* 32: 31-35.
2. Baldwin CL, Parent M (2002) Fundamentals of host immune response against *Brucella abortus*: what the mouse model has revealed about control of infection. *Veterinary Microbiology* 90: 367-382.
3. Ko J, Splitter GA (2003) Molecular host-pathogen interaction in brucellosis: current understanding and future approaches to vaccine development for mice and humans. *Clinical Microbiology Reviews* 16: 65-78.
4. Yagupsky P, Peled N, Press J, Abu-Rashid M, Abramson O (1997) Rapid detection of *Brucella melitensis* from blood cultures by a commercial system. *Eur J Clin Microbiol Infect Dis* 16: 605-607.
5. Shasha B, Lang R, Rubinstein E (1992) Therapy of experimental murine brucellosis with streptomycin, cotrimoxazole, ciprofloxacin, ofloxacin, pefloxacin, doxycycline, and rifampin. *Antimicrobial Agents and Chemotherapy* 36: 973-976.
6. Prior S, Gander B, Irache J M, Gamazo C (2005) Gentamicin loaded microspheres for treatment of experimental *Brucella abortus* infection in mice. *Journal of Antimicrobial Chemotherapy* 55: 1032-1036.
7. Izadjoo MJ, Mense MG, Bhattacharjee AK, Hadfield TL, Crawford RM, et al. (2008) A study on the use of male animal models for developing a live vaccine for brucellosis. *Transboundary and Emerging Diseases* 55: 145-151.
8. Shemesh AA, Yagupsky P (2011) Limitations of the standard agglutination test for detecting patients with *Brucella melitensis* bacteremia. *Vector Borne Zoonotic Dis* 11: 1599-1601.
9. McFarlane PA, Bayoumi AM (2004) Acceptance and rejection: cost-effectiveness and the working nephrologist. *Kidney International* 66: 1735-1741.
10. Okosun KO, Rachid O, Marcus N (2013) optimal control strategies and cost-effectiveness analysis of a malaria model. *Bio Systems* 111: 83-101.