

Stabilizing Anti-Tuberculosis Drugs with Methanol for Accurate Testing

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DESCRIPTION

Pharmacological testing is an essential step in evaluating the efficacy of anti-Tuberculosis (anti-TB) drugs, particularly in combating multidrug-resistant and extensively drug-resistant strains of *Mycobacterium tuberculosis* (*M. tb*). Preserving the activity and stability of these drugs during testing is essential for obtaining accurate results. Methanol, a commonly used organic solvent, has emerged as an effective agent for stabilizing anti-TB drugs in infected samples, ensuring reliability and reproducibility in pharmacological studies. The pharmacological assessment of anti-TB drugs involves testing their activity against live *M. tb* cultures or infected samples. Several challenges complicate this process, such as, drug degradation many anti-TB drugs are susceptible to degradation under laboratory conditions, leading to inaccurate efficacy data. Sample contamination biological samples often contain enzymes or reactive compounds that can degrade drugs. Maintaining drug potency by ensuring that the drugs retain their pharmacological activity throughout the testing period is important for meaningful results. Methanol has proven to be a reliable solution to address these challenges by preserving both the biological integrity of the samples and the chemical stability of the drugs.

Role and use of methanol in sample preservation

Methanol is widely used in microbiology and pharmacology for its unique properties, such as, fixative properties of methanol effectively fixes *M. tb* infected samples, halting microbial activity and preventing further metabolic degradation of drugs. Chemical stability provides a stable environment that minimizes drug degradation caused by oxidative or enzymatic processes. Methanol is compatible with various downstream analytical techniques, such as High-Performance Liquid Chromatography (HPLC), mass spectrometry, and spectrophotometry, ensuring accurate quantification of drug concentrations. In pharmacological testing, methanol is used to prepare and preserve *M. tb* infected samples as follows sample preparation infected samples are collected under sterile conditions and treated with methanol to inactivate the bacteria while preserving the sample's structural and chemical integrity. Anti-TB drugs are

introduced into the methanol-treated samples at desired concentrations. The samples are stored at controlled temperatures, ensuring long-term stability for extended testing periods. Preserved samples are subjected to pharmacological assays to evaluate drug efficacy, including Minimum Inhibitory Concentration (MIC) and bactericidal activity tests.

Applications in anti-tuberculosis drug research

Methanol-preserved samples are highly valuable in the development and evaluation of anti-TB drugs. Drug susceptibility testing of methanol-treated samples enable accurate determination of MIC values, guiding the selection of effective drug combinations. By stabilizing drugs in resistant *M. tb* strains, researchers can better understand resistance mechanisms and explore potential solutions. Methanol's compatibility with automated systems makes it ideal for large-scale screening of novel anti-TB compounds. Stable samples allow for precise measurements of drug activity over time, essential for optimizing dosing regimens. Moreover, methanol-preserved samples maintain the integrity of bacterial DNA and proteins, facilitating genomic and proteomic studies essential for identifying drug targets and resistance markers. This preservation method supports high-throughput workflows, ensuring reproducibility and consistency across experiments. Its application in combination with advanced analytical techniques enables detailed profiling of drug-bacteria interactions, accelerating the discovery of new therapeutic agents. Additionally, the ability to safely store and transport samples expands collaborative research opportunities, fostering advancements in global TB drug development initiatives.

Benefits of methanol preservation in TB research

Methanol's role in preserving *M. tb*-infected samples extends beyond pharmacological testing, offering broader benefits to TB research:

Enhanced safety: Methanol effectively inactivates live *M. tb*, reducing the risk of biosafety hazards during sample handling. This is particularly valuable in resource-limited settings where biosafety infrastructure may be inadequate.

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Facilitation of long term studies: Methanol-treated samples remain stable over extended periods, enabling longitudinal studies on drug efficacy and resistance trends without the risk of sample degradation.

Cost effective approach: Methanol is an affordable and widely available solvent, making it accessible for use in laboratories worldwide. Its dual role in inactivating bacteria and preserving drugs reduces the need for additional reagents or equipment, streamlining research workflows.

Support for collaborative research: Methanol-preserved samples can be safely transported between laboratories, fostering international collaborations. This facilitates multi-center studies, harmonizing research efforts to accelerate advancements in TB drug development.

Integration with omics technologies: Preserved samples retain the integrity of biomolecules such as DNA, RNA, and proteins.

This enables comprehensive genomic, transcriptomic, and proteomic analyses, providing insights into the molecular mechanisms of drug resistance and potential therapeutic targets.

CONCLUSION

The use of methanol to preserve *M. tb* infected samples represents a significant advancement in pharmacological testing of anti-TB drugs. By stabilizing drug potency and maintaining sample integrity, methanol enables researchers to generate accurate and reproducible data. This approach not only accelerates the development of new anti-TB therapies but also strengthens efforts to combat drug-resistant TB, ultimately contributing to improved global health outcomes.