





(IN202311005039)
An inhibitory compound against lumazine synthase and a method of preparing the same

Bacteria	Classification	Formula I	Formula I	Formula I
		Compound	Compound	Compound
		х=н	X=CI	X=CH <sub>3</sub>
Enterococcus	Gram +ve	6.25 μg/ ml	3.125 μg/ ml	3.125-6.25
faecium				μg/ ml
Streptococcus pneumoniae	Gram +ve	1.56 μg/ ml	1.56 μg/ ml	1.56 μg/ ml
Enterobacter	Gram -ve	50 μg/ ml	12.5-25 μg/	50 μg/ ml
sp.			ml	

### **NEED**

Drug-resistant tuberculosis (TB) remains a global health threat. Current treatments struggle against resistant strains, requiring new, more effective compounds.

# MARKET ANALYSIS The global tuberculosis dr

The global tuberculosis drugs market is projected to grow at a CAGR of 4.5%, reaching \$16.5 billion by 2033. Key growth drivers include rising TB cases, increased focus on antimicrobial resistance, and advancements in drug formulations.

## **TECHNOLOGY OVERVIEW**

This patent introduces an inhibitory compound targeting lumazine synthase, a key enzyme in Mycobacterium tuberculosis. When combined with rifampicin or isoniazid, it enhances the antibacterial activity against drug-resistant TB, offering a promising therapeutic solution.

## **Target Industries**

Biotechnology, Pharmaceuticals, Healthcare.

, Pharmaceutical manufacturers, biotechnology firms, healthcare providers, and research organizations focused on developing treatments for drug-resistant tuberculosis and other infectious diseases.

## **TECHNOLOGY KEY FEATURES**

A novel lumazine synthase inhibitor with enhanced efficacy against Mycobacterium tuberculosis when combined with rifampicin or isoniazid, reducing the minimum inhibitory concentration (MIC) for both compounds, showing potential for combating drug-resistant TB.

## AT A GLANCE

 SDG 3 (Good Health and Well-being), SDG 9 (Industry, Innovation, and Infrastructure)

#### Read more here

Technology is available for licensing/ co-development.

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