

## Ranbezolid

## Chemical Properties

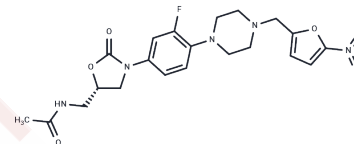
CAS No. : 392659-38-0

Formula: C<sub>21</sub>H<sub>24</sub>FN<sub>5</sub>O<sub>6</sub>

Molecular Weight: 461.44

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



## Biological Description

Description	Ranbezolid (RBx7644 free base) is an orally active oxazolidinone antibiotic effective against both Gram-positive and Gram-negative anaerobic bacteria, including <i>Staphylococcus aureus</i> , <i>Staphylococcus epidermidis</i> , and <i>Bacteroides fragilis</i> . It inhibits the 50s ribosomal subunit, with an IC <sub>50</sub> value of 17 μM for bacterial ribosomes, and disrupts cell wall and lipid synthesis. Ranbezolid acts as a rapid bactericidal agent, significantly reducing bacterial load, and demonstrates improved cardiovascular safety. It is applicable in the study of antibiotics targeting anaerobic bacteria.
Targets(IC50)	Antibacterial, Antibiotic
In vitro	Ranbezolid (1 μg/mL) effectively inhibits protein synthesis in <i>Staphylococcus aureus</i> and <i>Staphylococcus epidermidis</i> , while also impeding cell wall and lipid synthesis in <i>Staphylococcus epidermidis</i> . At concentrations between 1-8 μg/mL, Ranbezolid causes concentration-dependent membrane damage to <i>Staphylococcus epidermidis</i> . The compound's IC <sub>50</sub> for bacterial ribosomes is 17 μM, with a safety index of 865. Its minimum inhibitory concentrations (MIC) against the bacteria <i>Bacteroides fragilis</i> , <i>Bacteroides vulgatus</i> , <i>Bacteroides thetaiotaomicron</i> , <i>Clostridium perfringens</i> , <i>Clostridioides difficile</i> , and <i>Peptoniphilus magnus</i> are 0.06, 0.015, 0.06, 0.06, 0.03, and 0.015 μg/mL, respectively.
In vivo	Ranbezolid (100 mg/kg, p.o., administered twice daily for 5 days) demonstrates excellent in vivo activity against <i>Bacteroides fragilis</i> in a mouse infection model. In a rat spontaneous hypertension model, Ranbezolid (≤ 50 mg/kg, p.o., single dose) does not significantly enhance the pressor response induced by Tyramine.

### Preparing Stock Solutions

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	<b>1mg</b>	<b>5mg</b>	<b>10mg</b>
1 mM	2.1671 mL	10.8356 mL	21.6713 mL
5 mM	0.4334 mL	2.1671 mL	4.3343 mL
10 mM	0.2167 mL	1.0836 mL	2.1671 mL
50 mM	0.0433 mL	0.2167 mL	0.4334 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

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