

# BM635

Catalog No: tcsc0032638



## Available Sizes

**Size:** 1mg

**Size:** 5mg

**Size:** 10mg

**Size:** 25mg

**Size:** 50mg



## Specifications

**CAS No:**

1493762-74-5

**Formula:**

$C_{25}H_{29}FN_2O$

**Pathway:**

Anti-infection

**Target:**

Bacterial

**Purity / Grade:**

>98%

**Solubility:**

DMSO : 31 mg/mL (78.98 mM; Need ultrasonic and warming)

**Observed Molecular Weight:**

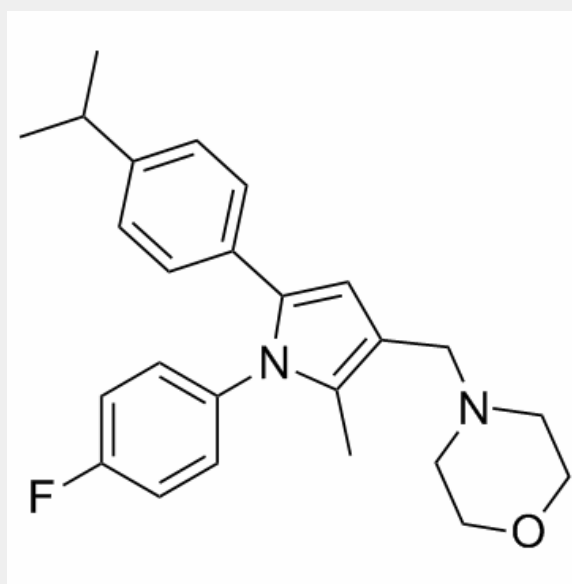
392.51

## Product Description

BM635 is a **MmpL3** inhibitor with outstanding anti-mycobacterial activity. BM635 has an **MIC<sub>50</sub>** of 0.12  $\mu$ M against *M. tuberculosis* H37Rv.

IC50 & Target: MIC50: 0.12  $\mu$ M (M. tuberculosis H37Rv)<sup>[1]</sup>

**In Vivo:** BM635 has potent MIC (0.12  $\mu$ M), Tox<sub>50</sub>:MIC ratio of >100, and good microsomal stability in mice (1.4 mL/min/g). When tested in an acute murine infection model at multiple doses, BM635 exhibits potent anti-tubercular activity, with an ED<sub>99</sub> of 49 mg/Kg (IC<sub>95%</sub>: 43–54 mg/Kg)<sup>[1]</sup>. The half-life *in vivo* of BM635 is 1h, allowing a reasonable maximum concentration (C<sub>max</sub>=1.62  $\mu$ M) and a moderate bioavailability (46%). Its poor aqueous solubility together with its high lipophilicity leads to low exposure *in vivo*<sup>[2]</sup>.



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!