## **Bio-AMS TFA**

Cat. No.:	HY-115448A		
Molecular Formula:	$C_{22}H_{30}F_{3}N_{9}O_{9}S_{2}$		
Molecular Weight:	685.65		
Target:	Bacterial		
Pathway:	Anti-infection		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month

Description	Bio-AMS (TFA) is a potent bacterial biotin protein ligase inhibitor. Bio-AMS (TFA) possesses selective activity against Mycobacterium tuberculosis (Mtb) and arrests fatty acid and lipid biosynthesis <sup>[1]</sup> .
IC <sub>50</sub> & Target	Biotin protein ligase <sup>[1]</sup>
In Vitro	Bio-AMS possesses excellent antitubercular activity against Mtb H37Rv and MDR/XDR-TB strains with MICs ranging from 0.16 to 0.625 μM and is not affected by changes to the primary carbon source <sup>[1]</sup> . Bio-AMS (2.5, 5 and 10 μM; 24 h) inhibits growth of Mtb in a concentration-dependent manner in Mtb-infected mouse macrophages; shows no signs of mitochondrial toxicity <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Bockman MR, Aldrich CC, et al. Avoiding Antibiotic Inactivation in Mycobacterium tuberculosis by Rv3406 through Strategic Nucleoside Modification. ACS Infect Dis. 2018 Jul 13;4(7):1102-1113.

[2]. Tiwari D, Schnappinger D, et al. Targeting protein biotinylation enhances tuberculosis chemotherapy. Sci Transl Med. 2018 Apr 25;10(438):eaal1803.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898 Fax: 609-228-5909 E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA



**BIOLOGICAL ACTIVITY** 

**Product** Data Sheet