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Data Sheet

Product Name:	Lalistat 1	0 <u> </u>
Cat. No.:	CS-0066587	
CAS No.:	501104-16-1	
Molecular Formula:	C ₁₂ H ₁₈ N ₄ O ₃ S	r
Molecular Weight:	298.36	O N
Target:	Bacterial	
Pathway:	Anti-infection	
Solubility:	DMSO : 20 mg/mL (67.03 mM; Need ultrasonic and warming)	

BIOLOGICAL ACTIVITY:

Lalistat 1 is a potent, selective, and competitive inhibitor of lysosomal acid lipase (LAL) and against purified human LAL (phLAL) with an IC50 of 68 nM. Lalistat 1 is a inhibitor of immunoglobulin A1 protease (IgA1P) proteases for H. influenzae, has less effects on other serine hydrolases (trypsin or β-lactamase, etc.). Lalistat 1 can be used for the research of niemann-pick type C (NPC) disease^[2]. IC50 & Target: IC50: 68 nM (human LAL)^[1] In Vitro: Lalistat 1 (0-100 µM) shows activity against different non-typeable H. influenzae (NTHi) IgAP variants from clinical isolates in Elisa assay. It shows a dose-dependent inhibition of IgA1P B1 and B2, but at higher inhibitor concentrations, consistent with the higher expression levels of these variants. Nearly complete inhibition of IgA1P B1 and B2 is observed at 50 μ M, with complete inhibition at 100 μ M^[1].

References:

[1]. Anton I Rosenbaum, et al. Thiadiazole carbamates: potent inhibitors of lysosomal acid lipase and potential Niemann-Pick type C disease therapeutics. J Med Chem . 2010 Jul 22;53(14):5281-9.

[2]. Livia Shehaj, et al. Small-Molecule Inhibitors of Haemophilus influenzae IgA1 Protease. ACS Infect Dis. 2019 Jul 12;5(7):1129-1138.

CAIndexNames:

4-Morpholinecarboxylic acid, 4-(1-piperidinyl)-1,2,5-thiadiazol-3-yl ester

SMILES:

O=C(N1CCOCC1)OC2=NSN=C2N3CCCCC3

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

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