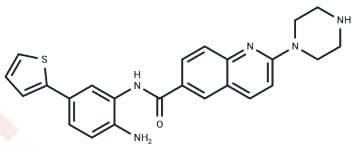
Data Sheet (Cat.No.T10245)

TargetMol

ACY-957

Chemical Properties



CAS No. : 1609389-52-7

Formula: C24H23N5OS

Molecular Weight: 429.54

Appearance: no data available

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

Biological Description

Description	ACY-957 (HDAC Inhibitor C001) is an orally active and selective inhibitor of HDAC1 and HDAC2 (IC50s: 7 nM, 18 nM, and 1300 nM against HDAC1/2/3) and shows no inhibition on HDAC4/5/6/7/8/9.
Targets(IC50)	HDAC
In vitro	ACY-957 is a selective inhibitor of HDAC1 and HDAC2, with IC50 values of 7 nM for HDAC1, 18 nM for HDAC2, and 1300 nM for HDAC3, showing no inhibition of HDAC4/5/6/7/8/9. It exhibits an IC50 of 304 nM for HDAC2 in primary hematopoietic progenitors [1].

Solubility Information

Solubility	DMSO: 50 mg/mL (116.4 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.3281 mL	11.6404 mL	23.2807 mL
5 mM	0.4656 mL	2.3281 mL	4.6561 mL
10 mM	0.2328 mL	1.164 mL	2.3281 mL
50 mM	0.0466 mL	0.2328 mL	0.4656 mL

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.

Reference

Shearstone JR, et al. Chemical Inhibition of Histone Deacetylases 1 and 2 Induces Fetal Hemoglobin through Activation of GATA2. PLoS One. 2016 Apr 13;11(4):e0153767.

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