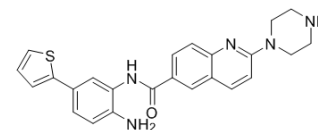


## ACY-957

Cat. No.:	HY-104008		
CAS No.:	1609389-52-7		
Molecular Formula:	C <sub>24</sub> H <sub>23</sub> N <sub>5</sub> OS		
Molecular Weight:	429.54		
Target:	HDAC		
Pathway:	Cell Cycle/DNA Damage; Epigenetics		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 83.33 mg/mL (194.00 mM; Need ultrasonic)				
	Preparing Stock Solutions	Solvent Concentration	1 mg	5 mg	10 mg
		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	1.1640 mL	2.3281 mL
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: <b>10% DMSO &gt;&gt; 40% PEG300 &gt;&gt; 5% Tween-80 &gt;&gt; 45% saline</b> Solubility: ≥ 2.08 mg/mL (4.84 mM); Clear solution				
	2. Add each solvent one by one: <b>10% DMSO &gt;&gt; 90% corn oil</b> Solubility: ≥ 2.08 mg/mL (4.84 mM); Clear solution				
	3. Add each solvent one by one: <b>10% DMSO &gt;&gt; 90% (20% SBE-β-CD in saline)</b> Solubility: ≥ 2.08 mg/mL (4.84 mM); Clear solution				

### BIOLOGICAL ACTIVITY

Description	ACY-957 is a selective inhibitor of HDAC1 and HDAC2, with IC <sub>50</sub> s of 7 nM, 18 nM, and 1300 nM against HDAC1/2/3, respectively, and shows no inhibition on HDAC4/5/6/7/8/9.		
IC <sub>50</sub> & Target	HDAC1 7 nM (IC <sub>50</sub> )	HDAC2 18 nM (IC <sub>50</sub> )	HDAC3 1300 nM (IC <sub>50</sub> )

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**In Vitro**

ACY-957 is a selective inhibitor of HDAC1 and HDAC2, with IC<sub>50</sub>s of 7 nM, 18 nM, and 1300 nM against HDAC1/2/3, respectively, and shows no inhibition on HDAC4/5/6/7/8/9. ACY-957 has an IC<sub>50</sub> of 304 nM for HDAC2 in primary hematopoietic progenitors<sup>[1]</sup>.

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**REFERENCES**

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

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