Data Sheet (Cat.No.T7057)



Methylstat

Chemical Properties

CAS No.: 1310877-95-2

Formula: C28H31N3O6

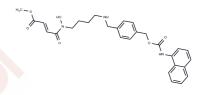
Molecular Weight: 505.56

Appearance: no data available

store at low temperature, keep away from direct

Storage: sunlight

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



Biological Description

Description	Methylstat is a methyl ester prodrug of a Jumonji C domain-containing histone		
	demethylase (JMJD) inhibitor that has favorable cell permeability. The free acid of		
	methylstat inhibits JMJD2A, JMJD2C, JMJD2E, PHF8, and JMJD3 with IC50 values of		
	approximately 4.3, 3.4, 5.9, 10, and 43 µM, respectively, in an in vitro assay. Methylstat		
	inhibits growth of the JMJD2C-sensitive esophageal cancer cell line KYSE150 with a GI50		
	of 5.1 μM, whereas the free acid of methylstat did not inhibit cell growth up to 100 μM.		
	Methylstat induces histone hypermethylation at multiple sites in a concentration-		
	dependent manner (EC50 for H3K4me3 and H3K9me3 = 10.3 and 8.6 µM in KYSE150 cells		
	and 6.7 ad 6.3 µM in MCF-7 cells, respectively).		
Targets(IC50)	Histone Demethylase		

Solubility Information

Solubility	DMSO: 25 mg/mL (49.45 mM),Sonication is recommended.
	(< 1 mg/ml refers to the product slightly soluble or insoluble)

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	1.978 mL	9.890 mL	19.780 mL
5 mM	0.3956 mL	1.978 mL	3.956 mL
10 mM	0.1978 mL	0.989 mL	1.978 mL
50 mM	0.0396 mL	0.1978 mL	0.3956 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.

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Reference

Luo, X., et al. A selective inhibitor and probe of the cellular functions of Jumonji C domain-containing histone demethylases. Journal of the American Chemical Society 133(24), 9451-9456 (2011).

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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