

Product Name: ML324 Revision Date: 10/14/2021

Product Data Sheet

ML324

Cat. No.: B4891

CAS No.: 1222800-79-4
Formula: C21H23N3O2

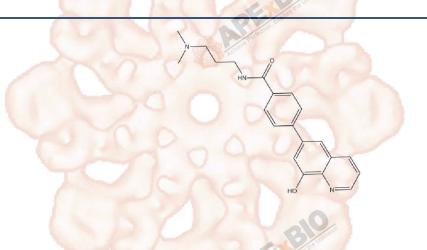
M.Wt: 349.43

Synonyms:

Target: Chromatin/Epigenetics

Pathway: Histone Demethylases

Storage: Store at -20°C



Solvent & Solubility

insoluble in H2O; \geq 17.45 mg/mL in DMSO; \geq 2.4 mg/mL in EtOH with ultrasonic

In Vitro

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	2.8618 mL	14.3090 mL	28.6180 mL
	5 mM	0.5724 mL	2.8618 mL	5.7236 mL
	10 mM	0.2862 mL	1.4 <mark>3</mark> 09 mL	2.8618 mL

Please refer to the solubility information to select the appropriate solvent.

Biological Activity

Shortsummary	JMJD2 demethylase inhibitor, potent and cell-permeable			
IC ₅₀ & Target		al ^o nom		
	Cell Viability Assay			
	Cell Line:	HFF cells		
	Preparation method:	The solubility of this compound in DMSO is > 17.5 mg/mL. General tips for		
In Vitro		obtaining a higher concentration: Please warm the tube at 37 °C for 10 minutes		
		and/or shake it in the ultrasonic bath for a while. Stock solution can be stored		
		below - 20 °C for several months.		
	Reacting conditions:	5 ~ 50 μM		

	Applications:	Compared with DMOG (IC50 = 0.75 mM), ML324 potently reduced IE gene		
		expression, with an IC50 value of 10 μ M. Besides, ML324 did not affect the		
		expression of the cellular controls Sp1, S15 and TBP. In HFF cells infected with		
		HSV-1, ML324 lowered viral yields in a dose-dependent manner (~ 4 ~ 5 logs at		
	The court	25 μM) while 1.5 mM of DMOG was required to cause the same reduction.		
	Animal experiment			
In Vivo	Animal models:	A mouse ganglia explant model of latently infected mice		
	Dosage form:	50 μM; 48 hrs		
	Applications:	In a mouse ganglia explant model of latently infected mice, ML324 significantly		
		inhibited viral activity. At the concentration of 50 µM, ML324 reduced the viral		
		yield by 4.5 logs for each ganglia. Immunofluorescent staining of explanted		
		ganglia sections showed that ML324 inhibited viral reactivation events.		
		However, the withdrawal of ML324 resulted in marked viral replication.		
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may		
	The Uthlicour	slightly differ with the theoretical value. This is caused by an experimental		
	California Explore	system error and it is normal.		

Product Citations

1. Li N, Yang L, et al. "BET bromodomain inhibitor JQ1 preferentially suppresses EBV-positive nasopharyngeal carcinoma cells partially through repressing c-Myc." Cell Death Dis. 2018 Jul 9;9(7):761.PMID:29988031

See more customer validations on www.apexbt.com.

References

[1]. Rai G, Kawamura A, Tumber A, Liang Y, Vogel JL, Arbuckle JH, Rose NR, Dexheimer TS, Foley TL, King ON, Quinn A, Mott BT, Schofield CJ, Oppermann U,Jadhav A, Simeonov A, Kristie TM, Maloney DJ. Discovery of ML324, a JMJD2 demethylase inhibitor with demonstrated antiviral activity. 2012 Dec 17 [updated 2013 Sep 16]. Probe Reports from the NIH Molecular Libraries Program [Internet]. Bethesda (MD): National Center for Biotechnology Information (US); 2010-.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.



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