**Proteins** 

## **Product** Data Sheet



Cat. No.: HY-19480 CAS No.: 548486-59-5 Molecular Formula:  $C_{12}H_{16}N_4O_3$ Molecular Weight: 264.28

Target: **Endogenous Metabolite** Pathway: Metabolic Enzyme/Protease -20°C Storage: Powder 3 years

4°C 2 years -80°C In solvent 6 months

> -20°C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 25 mg/mL (94.60 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.7839 mL	18.9193 mL	37.8387 mL
	5 mM	0.7568 mL	3.7839 mL	7.5677 mL
	10 mM	0.3784 mL	1.8919 mL	3.7839 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1 mg/mL (3.78 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1 mg/mL (3.78 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1 mg/mL (3.78 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description  $Ulodesine is a purine nucleoside phosphorylase (PNP) inhibitor. Ulodesine inhibits PNP with IC_{50} value of 2.293 \, nM/L. \\$ Ulodesine can be used for the research of hyporucicemia<sup>[1][2]</sup>. In Vivo

Ulodesine (i.v.) potently inhibits PNP with  $IC_{50}$  value of 2.293 nM/L<sup>[1]</sup>. Ulodesine (i.v.) eliminates uric acid accumulations in blood of the mouse  $model^{[1]}$ .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

]. Xujuan YANG, et al. Establis	hment of a novel hyperuricemiaanimal r	model using mice and ass	essment ofhyporuricemia action of PNP inhibitor Ulo	odesine.
]. Cesar Diaz-Torné, et al. Nev	v medications in development for the tre	eatment of hyperuricemia	of gout. Curr Opin Rheumatol. 2015 Mar;27(2):164-9.	
	Caution: Product has not been ful	lly validated for medica	al applications. For research use only.	
		509-228-5909	E-mail: tech@MedChemExpress.com	
	Address: 1 Deer Park	Dr, Suite Q, Monmouth	Junction, NJ 08852, USA	

**REFERENCES** 

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