Proteins

Inhibitors



M5N36

Cat. No.: HY-132578 CAS No.: 2832887-40-6 Molecular Formula: $C_{20}H_{16}CIN_5O_3$ Molecular Weight: 409.83

Target: Phosphatase

Pathway: Metabolic Enzyme/Protease Storage: Powder -20°C 3 years

> In solvent -80°C 6 months

-20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 25 mg/mL (61.00 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.4400 mL	12.2002 mL	24.4004 mL
	5 mM	0.4880 mL	2.4400 mL	4.8801 mL
	10 mM	0.2440 mL	1.2200 mL	2.4400 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: ≥ 2.5 mg/mL (6.10 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: ≥ 2.5 mg/mL (6.10 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	M5N36 is a potent and selective Cdc25C inhibitor with IC $_{50}$ values of 0.15, 0.19, 0.06 μ M for Cdc25A, Cdc25B, Cdc25C, respectively. M5N36 shows anti-proliferative activity and increases the expression of p-CDK1 and p-CDK2 $^{[1]}$.		
IC ₅₀ & Target	IC ₅₀ : 0.15 μM (Cdc25A); 0.19 μM (Cdc25B); 0.06 μM (Cdc25C) $^{[1]}$		
In Vitro	M5N36 (5, 10 μM; 24 h) increases the expression of p-CDK1 and p-CDK2 in MDA-MB-231 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay ^[1] Cell Line: Eca-109, HepG2, MCF-TAM, MDA-MB-231, Raji, SGC-7901, SW-620, HUVEC cells		

Concentration:		
Incubation Time:		
Result:	Showed anti-proliferative activity with IC $_{50}$ s of 2.88, 3.57, 10.00, 2.00, 1.20, 6.03, 3.39, 2.40 μ M for Eca-109, HepG2, MCF-TAM, MDA-MB-231, Raji, SGC-7901, SW-620, HUVEC cells, respectively.	
Western Blot Analysis ^[1]		
Cell Line:	MDA-MB-231 cells	
Concentration:	5, 10 μΜ	
Incubation Time:	24 h	
Result:	Increased the expression of Cdc25A, Cdc25B, increased the expression of p-CDK1 and p-CDK2.	

REFERENCES

[1]. Tao Y, et al. Discovery of potent and selective Cdc25 phosphatase inhibitors via rapid assembly and in situ screening of Quinonoid-focused libraries. Bioorg Chem. 2021 Oct;115:105254.

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

Tel: 609-228-6898

Fax: 609-228-5909

 $\hbox{E-mail: } tech@MedChemExpress.com$

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA