MNK1/2-IN-6

Cat. No.:	HY-146735	
CAS No.:	2771087-94-4	
Molecular Formula:	C ₂₇ H ₂₄ N ₆ O	
Molecular Weight:	448.52	N ^N
Target:	MNK; Apoptosis	H ₂ N H ₂ N
Pathway:	MAPK/ERK Pathway; Apoptosis	0 (= _N
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIV			
Description	MNK1/2-IN-6 is a potent and selective MNK1/2 inhibitor with IC ₅₀ s of 2.3 nM and 3.4 nM for MNK1 and MNK2, respectively. MNK1/2-IN-6 induces apoptosis in a concentration-dependent manner ^[1] .		
IC ₅₀ & Target	MNK1 2.3 nM (IC ₅₀)	MNK2 3.4 nM (IC ₅₀)	
In Vitro	MNK1/2-IN-6 (compound II-5) (72 h) shows antiproliferative activity with IC ₅₀ s of 0.3896μM, 0.4092 μM for TMD-8, DOHH-2 cells, respectively ^[1] . MNK1/2-IN-6 (0.25, 0.5, 1 μM; 24 h) induces TMD-8 cells apoptosis in a concentration-dependent manner ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay ^[1]		
	Cell Line:	TMD-8, DOHH-2 cells	
	Concentration:		
	Incubation Time:	72 h	
	Result:	Showed antiproliferative activity with $\rm IC_{50}s$ of 0.39 μ M, 0.41 μ M for TMD-8, DOHH-2 cells, respectively.	
	Apoptosis Analysis ^[1]		
	Cell Line:	TMD-8 cells	
	Concentration:	0.25, 0.5, 1 μΜ	
	Incubation Time:	24 h	
	Result:	Induced TMD-8 cells apoptosis in a concentration-dependent manner.	

REFERENCES

Product Data Sheet



[1]. Bu H, et al. Design, synthesis and biological evaluation of imidazopyridazine derivatives containing isoquinoline group as potent MNK1/2 inhibitors. Bioorg Med Chem. 2021; 40:116186.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

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