Product Data Sheet

Ubiquitination-IN-2

Cat. No.: HY-152266 Molecular Formula:

Molecular Weight: 2346.04

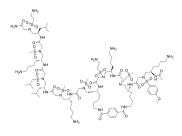
Target: E1/E2/E3 Enzyme

Pathway: Metabolic Enzyme/Protease

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

 $\mathsf{C_{93}H_{173}N_{25}O_{28}S_8}$



BIOLOGICAL ACTIVITY

Description	Ubiquitination-IN-2 is a potent E1-E2 protein–protein interactions (PPI) inhibitor. Ubiquitination-IN-2 has a K_d value of 0.72 μ M for ubiquitin E1 (Uba1). Ubiquitination-IN-2 inhibits blocks ubiquitin transfer from E1 to E2. Ubiquitination-IN-2 can be used in research of cancer ^[1] .	
In Vitro	Ubiquitination-IN-2 (M1-S1; 0-50 μ M; 16 h; HEK293T cells) inhibits ubiquitin transfer to substrate protein CDK4 by blocking its monoubiquitination catalyzed by CHIP ^[1] . Ubiquitination-IN-2 (0-50 μ M; HEK293T cells) inhibits the protein ubiquitination cascade by binding with Uba1, the E1 enzyme, and blocking ubiquitin transfer from E1 to E2 ^[1] . Ubiquitination-IN-2 (0-50 μ M; HEK293T cells) inhibits ubiquitin transfer to different types of E2 enzymes and inhibits the formation of E2-UB conjugates at 50 μ M concentration ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis ^[1]	
	Cell Line:	HEK293T cells
	Concentration:	0, 10, 50 μΜ
	Incubation Time:	16 hours
	Result:	Inhibitd the ubiquitination of CDK4 in HEK293T cells.

REFERENCES

[1]. Zhou L, et, al. Inhibition of the Ubiquitin Transfer Cascade by a Peptidomimetic Foldamer Mimicking the E2 N-Terminal Helix. J Med Chem. 2023 Jan 12;66(1):491-502.

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