CBP/p300-IN-19

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Cat. No.:	HY-146277	
CAS No.:	2592638-13-4	
Molecular Formula:	$C_{30}H_{27}N_{3}O_{3}$	
Molecular Weight:	477.55	N
Target:	Histone Acetyltransferase	
Pathway:	Epigenetics	HN
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	~0

BIOLOGICAL ACTI	VITY				
Description	CBP/p300-IN-19 is a potent p300/CBP HAT inhibitor with IC ₅₀ s of 1.4, 2.2, >100, >100 μM for p300-HAT, CBP-HAT, PCAF, Myst3, respectively. CBP/p300-IN-19 shows antitumor activity ^[1] .				
IC ₅₀ & Target	Myst3 >100 μM (IC ₅₀)	р300-НАТ 1.4 µМ (IC ₅₀)	CBP-HAT 2.2 μΜ (IC ₅₀)	PCAF >100 μM (IC ₅₀)	
In Vitro	CBP/p300-IN-19 (compound 29) shows antiproliferative activity with EC ₅₀ s of 5.3, 8.5, 6.2, 4.4, 1.2, 4.3, 3.6, 8.7, 6.4 μM for MCF-7, MDA-MB231, LNCaP, PC-3, PANC-1, MDA-PANC-28, Molm-13, MV4;11, RPMI-8226 cells, respectively ^[1] . CBP/p300-IN-19 (0, 5, 10 μM; 12 h) dose-dependenly inhibits the acetylation of H3K18, H3K9 and K27 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis ^[1]				
	Cell Line:	Kasumi-1 leukemia cells			
	Concentration:	0, 5, 10 μΜ			
	Incubation Time:	12 h			
	Result:	Dose-dependently inhibited the acetylation of H3K18, H3K9 and K27.			

REFERENCES

[1]. Nie S, et al. Structure-activity relationship and antitumor activity of 1,4-pyrazine-containing inhibitors of histone acetyltransferases P300/CBP. Eur J Med Chem. 2022 Jul 5;237:114407.

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

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