Suramin

Cat. No.: CAS No.:	HY-B0879 145-63-1	
Molecular Formula:	C ₅₁ H ₄₀ N ₆ O ₂₃ S ₆	"Å~ • • • ~ • ~ • • • •
Molecular Weight: Target:	1297.28 Phosphatase; Sirtuin; Reverse Transcriptase; Topoisomerase; Apoptosis; Parasite; SARS-CoV	
Pathway:	Metabolic Enzyme/Protease; Cell Cycle/DNA Damage; Epigenetics; Anti-infection; Apoptosis	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIV				
Description	Suramin is a reversible and competitive protein-tyrosine phosphatases (PTPases) inhibitor ^[1] . Suramin is a potent inhibitor of sirtuins: SirT1 (IC ₅₀ =297 nM), SirT2 (IC ₅₀ =1.15 μ M), and SirT5 (IC ₅₀ =22 μ M) ^[2] . Suramin is a competitive inhibitor of reverse transcriptase (DNA topoisomerase II: IC ₅₀ =5 μ M) ^{[3][4]} . Suramin is a potent SARS-CoV-2 RNA-dependent RNA polymerase (RdRp) inhibitor ^[5] . Suramin efficiently inhibits IP5K and is an antiparasitic, anti-neoplastic and anti-angiogenic agent ^{[6][7][8]} .			
IC ₅₀ & Target	SIRT1 297 nM (IC ₅₀)	SIRT2 1.15 μΜ (IC ₅₀)	SIRT5 22 μΜ (IC ₅₀)	
In Vitro	Suramin (50-600 μg/mL; for 24-96 hours) inhibits cells proliferation in a dose-dependent and time-dependent manner and decreases viability in cancer cells ^[7] . Suramin (300 μg/mL; for 48 hours) induces cells apoptosis, and down-regulates mRNA expression in HeLa cells ^[7] . Suramin (1 mg/mL; 1 hour) significantly suppresses the phosphorylated ERK1/2 ^[8] . The IC ₅₀ values of HO-8910 PM and HeLa are 319 μg/mL, 476 μg/mL, respectively ^[7] . Suramin blocks viral replication in Vero E6 cells ^[5] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay ^[6]			
	Cell Line:	HO-8910 PM ovarian and Hela	cervical cancer cells	
	Concentration:	50, 100, 200, 300, 400, 500 and 600 μg/mL		
	Incubation Time:	For 24, 48, 72 and 96 hours		
	Result:	Inhibited cells proliferation in a dose-dependent and time-dependent manner.		
	Apoptosis Analysis ^[6]			
	Cell Line:	HeLa cells		
	Concentration:	300 μg/mL		
	Incubation Time:	For 48 hours		

Product Data Sheet



	Result:	Induced cells apoptosis.		
	Western Blot Analysis ^[7]	: 		
	Cell Line:	PA-SMCs cells		
	Concentration:	1 mg/mL		
	Incubation Time:	For 1 hours		
	Result:	Significantly suppressed the phosphorylated ERK1/2.		
In Vivo	the pulmonary artery p	Suramin (10 mg/kg; IV; twice weekly for 3 weeks) reverses established pulmonary hypertension (PH), thereby normalizing the pulmonary artery pressure values and vessel structure ^[8] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Adult male Wistar rats (200-225 g) ^[7]		
	Dosage:	10 mg/kg		
	Administration:	IV; twice weekly for 3 weeks		
	Result:	Reversed established PH, thereby normalizing the pulmonary artery pressure values and vessel structure.		

CUSTOMER VALIDATION

- Nat Struct Mol Biol. 2021 Mar;28(3):319-325.
- Clin Transl Med. 2021 Jun;11(6):e485.
- Br J Pharmacol. 2021 Aug 6.
- J Agric Food Chem. 2023 Sep 19.
- Int Immunopharmacol. 2023 May 12;120:110295.

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[1]. Jindal HK, et al. Suramin affects DNA synthesis in HeLa cells by inhibition of DNA polymerases. Cancer Res. 1990 Dec 15;50(24):7754-7.

[2]. Izikki M, et al. The beneficial effect of suramin on monocrotaline-induced pulmonary hypertension in rats. PLoS One. 2013 Oct 15;8(10):e77073.

[3]. Zhang YL, et al. Suramin is an active site-directed, reversible, and tight-binding inhibitor of protein-tyrosine phosphatases. J Biol Chem. 1998 May 15;273(20):12281-7.

[4]. Trapp J, et al. Structure-activity studies on suramin analogues as inhibitors of NAD+-dependent histone deacetylases (sirtuins). ChemMedChem. 2007 Oct;2(10):1419-31.

[5]. Schuetz A, et al. Structural basis of inhibition of the human NAD⁺-dependent deacetylase SIRT5 by suramin. Structure. 2007 Mar;15(3):377-89.

[6]. De Clercq E, et al. Suramin: a potent inhibitor of the reverse transcriptase of RNA tumor viruses. Cancer Lett. 1979 Nov;8(1):9-22.

[7]. Novaes RD, et al. Purinergic Antagonist Suramin Aggravates Myocarditis and Increases Mortality by EnhancingParasitism, Inflammation, and Reactive Tissue Damage in Trypanosoma cruzi-Infected Mice. Oxid Med Cell Longev. 2018 Sep 30;2018:7385639. [8]. Xiaozhe Zhang, et al. Suramin and NF449 Are IP5K Inhibitors That Disrupt IP6-mediated Regulation of Cullin RING Ligase and Sensitize Cancer Cells to MLN4924/pevonedistat. J Biol Chem. 2020 Jun 3; jbc.RA120.014375.

[9]. Wanchao Yin, et al. Structural basis for inhibition of the SARS-CoV-2 RNA polymerase by suramin. Nat Struct Mol Biol. 2021 Mar;28(3):319-325.

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

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