Geldanamycin

Cat. No.:	HY-15230			
CAS No.:	30562-34-6			
Molecular Formula:	$C_{29}H_{40}N_2O_9$			
Molecular Weight:	560.64			
Target:	HSP; Bacterial; Influenza Virus; Antibiotic			
Pathway:	Cell Cycle/DNA Damage; Metabolic Enzyme/Protease; Anti-infection			
Storage:	Powder	-20°C	3 years	
		4°C	2 years	
	In solvent	-80°C	6 months	
		-20°C	1 month	

SOLVENT & SOLUBILITY

In Vitro	DMSO : 25 mg/mL (44.59 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	1.7837 mL	8.9184 mL	17.8368 mL	
		5 mM	0.3567 mL	1.7837 mL	3.5674 mL	
		10 mM	0.1784 mL	0.8918 mL	1.7837 mL	
	Please refer to the solubility information to select the appropriate solvent.					
In Vivo	 Add each solvent of Solubility: 2.5 mg/ Add each solvent of Solubility: 2.5 mg/ 	one by one: 10% DMSO >> 40% PEC mL (4.46 mM); Suspended solution; one by one: 10% DMSO >> 90% (20 mL (4.46 mM); Suspended solution;	5300 >> 5% Tween-80 Need ultrasonic % SBE-β-CD in saline) Need ultrasonic) >> 45% saline		

biological activity				
Description	Geldanamycin is a Hsp90 inhibitor with antimicrobial activity against many Gram-positive and some Gram-negative bacteria. Geldanamycin has anti-influenza virus H5N1 activities.			
IC ₅₀ & Target	HSP90 1.2 μM (Kd)			
In Vitro	Geldanamycin significantly delays and reduces viperin expression, indicating that IRF3 is involved in viperin induction in RAW264.7 cells ^[1] . Geldanamycin, a benzoquinone ansamycin, protected against neuronal injury induced by oxygen-glucose deprivation (OGD)/zVAD treatment in cultured primary neurons. More importantly, Geldanamycin decreases RIP1 protein level in a time			

OH

0:

Ô

ΝH₂

O

0



and concentration-dependent manner. Geldanamycin also decreases the Hsp90 protein level, which causes instability of RIP1 protein, resulting in decreased RIP1 protein level but not RIP1 mRNA level after Geldanamycin treatment^[2]. Geldanamycin is identified as the first natural product inhibitor of Hsp90 that binds to the N-terminal ATPase domain of Hsp90 to inhibit its chaperone function, and significantly induces tumor cell death via an apoptotic mechanism^[3]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Nat Commun. 2022 Aug 29;13(1):4942.
- Nat Commun. 2021 May 10;12(1):2587.
- Nucleic Acids Res. 2020 Aug 20;48(14):7944-7957.
- Autophagy. 2023 Jun 13;1-17.
- New Phytol. 2023 Jan 27.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Tang HB, et al. Viperin inhibits rabies virus replication via reduced cholesterol and sphingomyelin and is regulated upstream by TLR4. Sci Rep. 2016 Jul 26;6:30529

[2]. Chen WW, et al. RIP1 mediates the protection of Geldanamycin on neuronal injury induced by oxygen-glucosedeprivation combined with zVAD in primary cortical neurons. J Neurochem. 2012 Jan;120(1):70-7.

[3]. Lin Z, et al. 17-ABAG, a novel Geldanamycin derivative, inhibits LNCaP-cell proliferation through heat shock protein 90 inhibition. Int J Mol Med. 2015 Aug;36(2):424-32.

[4]. Roe SM, et al. Structural basis for inhibition of the Hsp90 molecular chaperone by the antitumor antibiotics radicicol and geldanamycin. J Med Chem. 1999 Jan 28;42(2):260-6.

[5]. Wang C, et al. Geldanamycin Reduces Acute Respiratory Distress Syndrome and Promotes the Survival of Mice Infected with the Highly Virulent H5N1 Influenza Virus. Front Cell Infect Microbiol. 2017 Jun 15;7:267.

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