Ganoderic acid D

Cat. No.:	HY-N1511			
CAS No.:	108340-60-9			
Molecular Formula:	C ₃₀ H ₄₂ O ₇			
Molecular Weight:	514.65			
Target:	Sirtuin; Apo	ptosis		
Pathway:	Cell Cycle/DNA Damage; Epigenetics; Apoptosis			
Storage:	Powder	-20°C	3 years	
		4°C	2 years	
	In solvent	-80°C	6 months	
		-20°C	1 month	

SOLVENT & SOLUBILITY

	Preparing Stock Solutions	Solvent	1 mg	5 mg	10 mg		
		Concentration	8	g			
		1 mM	1.9431 mL	9.7153 mL	19.4307 mL		
		5 mM	0.3886 mL	1.9431 mL	3.8861 mL		
		10 mM	0.1943 mL	0.9715 mL	1.9431 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (4.86 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.86 mM); Clear solution						
	3. Add each solvent	 Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.86 mM); Clear solution 					

BIOLOGICAL ACTIVITY

Description	Ganoderic acid D, a highly oxygenated tetracyclic triterpenoid, is the major active component of Ganoderma lucidum.
	Ganoderic acid D upregulates the protein expression of SIRT3 and induces the deacetylated cyclophilin D (CypD) by SIRT3.
	Ganoderic acid D inhibits the energy reprogramming of colon cancer cells including glucose uptake, lactate production,
	pyruvate and acetyl-coenzyme production in colon cancer cells ^[1] . Ganoderic acid D induces HeLa human cervical carcinoma
	apoptosis ^[2] .

Product Data Sheet

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n Vitro	with an IC ₅₀ of 17.3 mM [[] dependent manner ^[2] . Ganoderic acid D (10, 50 Ganoderic acid D (10, 50 Ganoderic acid D (10 µM	hibit the growth of numerous cancer cell lines and it inhibits HeLa human cervical carcinoma cells ^{2]} .Ganoderic acid D (1-50 μM; 24-72 hours) reduces the cell survival rate in a dose- and time- μμM; 24, 48 hours) induces G2/M phase arrest ^[2] . μμM; 24, 48 hours) induces a morphological change typical of apoptosis in HeLa cells ^[2] . I; 48 hours) up-regulates 14-3-3E and PRDX3 ^[2] . htly confirmed the accuracy of these methods. They are for reference only.			
	Cell Line:	HeLa human cervical carcinoma cell line (CCL-2)			
	Concentration:	1, 5, 10, 20, 50 μM			
	Incubation Time:	24, 48, 72 hours			
	Result:	Reduced the cell survival rate in a dose- and time-dependent manner and had an IC_{50} value of 17.3 μM for 48 hours treatment.			
	Cell Cycle Analysis ^[2]				
	Cell Line:	HeLa human cervical carcinoma cell line (CCL-2)			
	Concentration:	10, 50 μM			
	Incubation Time:	24, 48 hours			
	Result:	Induced G2/M phase arrest. Displayed a cell cycle profile with an elevated G2/M cell population after 24-h treatment with 10 $\mu M.$			
	Apoptosis Analysis ^[2]				
	Cell Line:	HeLa human cervical carcinoma cell line (CCL-2)			
	Concentration:	10, 50 μM			
	Incubation Time:	48 hours			
	Result:	Induced a morphological change typical of apoptosis in HeLa cells.			
	Western Blot Analysis ^[2]	Western Blot Analysis ^[2]			
	Cell Line:	HeLa human cervical carcinoma cell line (CCL-2)			
	Concentration:	10 μΜ			
	Incubation Time:	48 hours			
	Result:	Up-regulated 14-3-3E and PRDX3.			

CUSTOMER VALIDATION

• Front Pharmacol. 2022 Feb 21:13:826716.

REFERENCES

[1]. Liu Z, et al. Effect of ganoderic acid D on colon cancer Warburg effect: Role of SIRT3/cyclophilin D. Eur J Pharmacol. 2018 Apr 5;824:72-77.

[2]. Yue QX, et al. Proteomics characterization of the cytotoxicity mechanism of ganoderic acid D and computer-automated estimation of the possible drug target network. Mol Cell Proteomics. 2008 May;7(5):949-61.

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

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