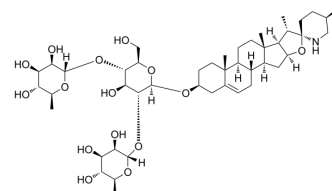


Solamargine

Cat. No.:	HY-N0069		
CAS No.:	20311-51-7		
Molecular Formula:	C ₄₅ H ₇₃ NO ₁₅		
Molecular Weight:	868.06		
Target:	P-glycoprotein; Apoptosis; P-glycoprotein		
Pathway:	Membrane Transporter/Ion Channel; Apoptosis		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (115.20 mM; Need ultrasonic)					
		Solvent Concentration	Mass	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	1.1520 mL	5.7600 mL	11.5199 mL	
		5 mM	0.2304 mL	1.1520 mL	2.3040 mL	
10 mM		0.1152 mL	0.5760 mL	1.1520 mL		
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (2.88 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (2.88 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (2.88 mM); Clear solution 					

BIOLOGICAL ACTIVITY

Description	Solamargine, a derivative from the steroidal solasodine in Solanum species, exhibits anticancer activities in numerous types of cancer. Solamargine induces non-selective cytotoxicity and P-glycoprotein inhibition. Solamargine significantly inhibits migration and invasion of HepG2 cells by down-regulating MMP-2 and MMP-9 expression and activity ^{[1][2]} .
In Vitro	Solamargine (15 µg/ml; 72 hours) predominantly stimulates the cells of the G2/M phase to apoptosis ^[3] . Solamargine (72 hours) dramatically inhibits the proliferation of SMMC-7721 and HepG2 cells in a dose- and time-dependent manner. The IC ₅₀ s are 9.21 and 19.88 µg/ml, respectively ^[3] .

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay^[3]

Cell Line:	SMMC-7721cells
Concentration:	15 µg/ml
Incubation Time:	72 hours
Result:	Displayed a significant increase of sub-G1.

In Vivo

Solamargine (10 mg/kg; intragastric administration; once daily for 8 days) exhibits an antitumor effect and promotes the apoptosis of GC in vivo^[4].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Female specific pathogen-free BALB/c nude mice weighing 18-20 g (6-8-weeks-old) ^[4]
Dosage:	10 mg/kg
Administration:	Intragastric administration; once daily for 8 days
Result:	Tumor growth was significantly inhibited .

CUSTOMER VALIDATION

- Food Chem. 2020 May 15;312:126030.
- Cancers. 2019 Mar 12;11(3):353.
- Int J Oncol. 2019 Mar;54(3):905-915.
- Int J Oncol. 2019 May;54(5):1545-1554.
- J Sci Food Agric. 2019 May;99(7):3578-3587.

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REFERENCES

- [1]. Burger T, et al. Solamargine, a bioactive steroidal alkaloid isolated from *Solanum aculeastrum* induces non-selective cytotoxicity and P-glycoprotein inhibition. *BMC Complement Altern Med*. 2018 May 2;18(1):137.
- [2]. Sani IK, et al. Solamargine inhibits migration and invasion of human hepatocellular carcinoma cells through down-regulation of matrix metalloproteinases 2 and 9 expression and activity. *Toxicol In Vitro*. 2015 Aug;29(5):893-900.
- [3]. Ding X, et al. Induction of apoptosis in human hepatoma SMMC-7721 cells by solamargine from *Solanum nigrum* L. *J Ethnopharmacol*. 2012 Jan 31;139(2):599-604.
- [4]. Fu R, et al. Solamargine inhibits gastric cancer progression by regulating the expression of *lncNEAT1_2* via the MAPK signaling pathway. *Int J Oncol*. 2019 May;54(5):1545-1554.

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

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