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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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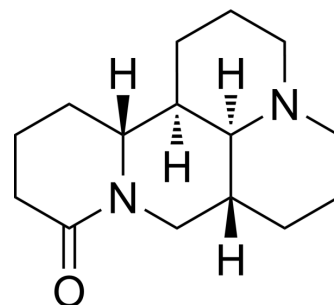
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Sophoridine

Cat. No.:	HY-N1373		
CAS No.:	6882-68-4		
Molecular Formula:	C ₁₅ H ₂₄ N ₂ O		
Molecular Weight:	248.36		
Target:	Apoptosis		
Pathway:	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 : 50 mg/mL (201.32 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	4.0264 mL	20.1321 mL	40.2641 mL
		5 mM	0.8053 mL	4.0264 mL	8.0528 mL
10 mM		0.4026 mL	2.0132 mL	4.0264 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: ≥ 2.17 mg/mL (8.74 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.17 mg/mL (8.74 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.17 mg/mL (8.74 mM); Clear solution 				

BIOLOGICAL ACTIVITY

Description	Sophoridine is a quinolizidine alkaloid isolated from Leguminous plant <i>Sophora flavescens</i> . Sophoridine induces apoptosis. Sophoridine has the potential to be a novel, potent and selective antitumor agent candidate for pancreatic cancer with well-tolerated toxicity ^[1] .
In Vitro	Sophoridine (0-500 μM; 48 hours) exhibits remarkable inhibition effects to the growth of human pancreatic, gastric, liver, colon, gallbladder, and prostate carcinoma cells with IC ₅₀ values of about 20 μM to 200 μM ^[1] . Sophoridine (0-20 μM; 48 hours) increases S phase cell population from 26.23% (control) to 38.67% in Miapaca-2 cells and

from 29.56% (control) to 39.16% in PANC-1 cells, about a 1.5-fold and a 1.3-fold increase, respectively^[1]. Sophoridine (0-20 μ M; 48 hours) significantly increases bad and bax levels, and decreases bcl-2 and bcl-xl levels in contrast, with a significant increase in Bax/Bcl-2 ratio^[1].

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

Cell Viability Assay^[1]

Cell Line:	Normal cells: IOSE144, HL-7702 and LO2, BEAS-2B, GES-1, HEK 293 T, HPDE, FHC, Human cancer cells: PANC-1, Mapaca-1, hepG2, SGC-7901, CBC-SD,SGC-996,PC-3,MKN-45,MGC-803,Hela and HCT116 cells
Concentration:	0, 3.9, 7.8, 15.5, 31, 62.5, 125, 250, 500 μ M
Incubation Time:	48 hours
Result:	Exhibited the most potent cytotoxicity to cancer cells.

Cell Cycle Analysis^[1]

Cell Line:	PANC-1 cells; Miapca-2 cells
Concentration:	20 μ M
Incubation Time:	48 hours
Result:	Led to accumulated population in the S phase.

Western Blot Analysis^[1]

Cell Line:	PANC-1 cells; Miapca-2 cells
Concentration:	20 μ M
Incubation Time:	48 hours
Result:	Induced the activation of intrinsic apoptosis pathway.

In Vivo

Sophoridine (intraperitoneal injection; 20 or 40 mg/kg; 21 days) can inhibit the growth of xenograft pancreatic tumors^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	BALB/c homozygous (nu/nu) nude mice ^[1]
Dosage:	20 or 40 mg/kg
Administration:	Intraperitoneal injection; 20 or 40 mg/kg; 21 days
Result:	Decreased xenograft pancreatic tumors mass.

CUSTOMER VALIDATION

- Front Oncol. 2021 Feb 24;11:634851.
- J Food Nutr Res. 2021, 9(11), 591-596.
- Research Square Preprint. 2020 Jun.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Xu Z, et al. Sophoridine induces apoptosis and S phase arrest via ROS-dependent JNK and ERK activation in human pancreatic cancer cells. J Exp Clin Cancer Res. 2017 Sep 11;36(1):124.

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

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