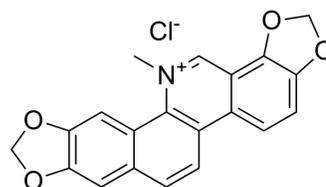


Sanguinarine chloride

Cat. No.:	HY-N0052A
CAS No.:	5578-73-4
Molecular Formula:	C ₂₀ H ₁₄ ClNO ₄
Molecular Weight:	367.78
Target:	Apoptosis; Autophagy; Bacterial; Parasite
Pathway:	Apoptosis; Autophagy; Anti-infection
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 1 year; -20°C, 6 months (sealed storage, away from moisture and light)



SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 6 mg/mL (16.31 mM; Need ultrasonic)				
	DMSO : 3.33 mg/mL (9.05 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.7190 mL	13.5951 mL	27.1902 mL
	5 mM	0.5438 mL	2.7190 mL	5.4380 mL	
	10 mM	0.2719 mL	1.3595 mL	2.7190 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: ≥ 0.33 mg/mL (0.90 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 0.33 mg/mL (0.90 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Sanguinarine (Sanguinarin) chloride, a benzophenanthridine alkaloid derived from the root of Sanguinaria Canadensis, can stimulate apoptosis via activating the production of reactive oxygen species (ROS). Sanguinarine-induced apoptosis is associated with the activation of JNK and NF-κB.
IC₅₀ & Target	Apoptosis ^[1]
In Vitro	Sanguinarine (SANG)-induced apoptosis is associated with the activation of JNK and NF-κB signal pathways. To determine the effects of Sanguinarine on cell viability, 22B-cFluc cells are stimulated with different concentrations of Sanguinarine for 24 h, and then a CKK-8 assay is performed. The treatment with Sanguinarine decreases the proliferation of 22B cells in a dose-dependent manner. Meanwhile, the cytosolic extracts of 22B-cFluc cells treated with different dose of Sanguinarine are

measured to detect cellular caspase-3 activity using Ac-DEVD-pNA, which is a validated caspase-3 substrate. The absorbance at 450 nm increases in a dose-dependent manner, indicating increased caspase-3 activity stimulated by Sanguinarine^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

In Vivo

To evaluate the apoptosis induced by Sanguinarine (SANG) in vivo, 22B-cFluc cells are inoculated subcutaneously into one flank of nude mice and xenograft models are allowed to establish. Mice are treated intravenously with 10 mg/kg of Sanguinarine. At 24, 48 and 72 h after treatment, bioluminescent imaging is performed after i.p. injection of mice with 150 mg/kg of D-luciferin substrate. Sanguinarine treatment induces an obvious increase of luminescent signal as early as 48 h after initial treatment. A sustained bioluminescent imaging (BLI) intensity increased is observed throughout the course of experiment. At 72 h after treatment, the tumors are collected and subjected to TUNEL staining for evaluating apoptosis. Compared with the control tumors, the group treated with Sanguinarine exhibits significantly more cell apoptosis, indicated by the increased green signals from the sporadic apoptotic cells^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Kinase Assay ^[1]

The caspase-3 activity is measured using a caspase-3 activity assay kit. Briefly, the cells treated by different concentrations of Sanguinarine (0.5 μ M, 1 μ M, 2 μ M, 4 μ M) or control DMSO are collected, washed and lysed in a lysis buffer for 30 min on ice. The supernatants are then collected by centrifuging at 1,2000 rpm for 10 min. The Ac-DEVD-pNA (2 mM) is added to each sample and incubated at 37°C for 1 h. The optical density (OD) of each sample is finally quantified at a wavelength of 405 nm using a spectrophotometer. The p-NA standard is used to calibrate the caspase-3 activity of each sample^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Assay ^[1]

The cell viability of Sanguinarine is determined by CCK-8 assay using a cell counting kit-8. Briefly, 22B-cFluc cells are seeded in a 96-well plate (5×10^3 cells/well) and treated with different concentrations of Sanguinarine (0.5 μ M, 1 μ M, 2 μ M, 4 μ M) for 24 h. Then 10 mL CCK-8 is added to each well for 4 h and the absorbance at 450 nm is measured with a microplate reader. The optical density (OD) values are determined to reflect the viable cell populations from each well^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Administration ^[1]

Mice^[1]
Xenografted tumor models are prepared by injection of 1×10^7 22B-cFluc cells suspended in PBS into nude mouse (n=6). After tumors reach a volume of approximately 100 mm³, Sanguinarine (10 mg/kg) is i.v. injected into mice. After injection for 24 h, 48 h and 72 h, mice are given a single i.p. dose of 150 mg/kg D-luciferin and bioluminescence imaging are performed using a Xenogen Lumina II system. The signal intensity in the region of interest is expressed using the Living Image software 4.1. For the anti-tumor therapy studies, one group of tumor-bearing mice (n=6) receive intravenously 10 mg/kg of Sanguinarine every other day throughout the experimental period, while the control group of mice (n=6) receive DMSO only. Tumor growth measurement is calculated^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Phytother Res. 2023 Jul 11.
- PLoS Pathog. 2023 Dec 7;19(12):e1011796.
- Pestic Biochem Physiol. 10 October 2022, 105259.
- Comp Biochem Physiol C Toxicol Pharmacol. 28 October 2021, 109228.
- Toxicol Lett. 2021 Jul 9;S0378-4274(21)00180-6.

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REFERENCES

[1]. Wang Y, Noninvasive bioluminescence imaging of the dynamics of sanguinarine induced apoptosis via activation of reactive oxygen species. Oncotarget. 2016 Apr 19;7(16):22355-67.

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

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