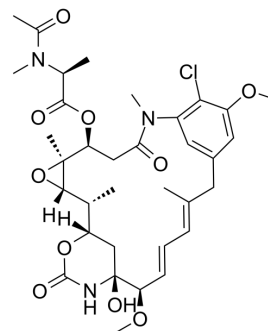


Maytansine

Cat. No.:	HY-13674		
CAS No.:	35846-53-8		
Molecular Formula:	C ₃₄ H ₄₆ ClN ₃ O ₁₀		
Molecular Weight:	692		
Target:	Microtubule/Tubulin		
Pathway:	Cell Cycle/DNA Damage; Cytoskeleton		
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 : 100 mg/mL (144.51 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.4451 mL	7.2254 mL	14.4509 mL
5 mM	0.2890 mL	1.4451 mL	2.8902 mL
10 mM	0.1445 mL	0.7225 mL	1.4451 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.5 mg/mL (3.61 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.5 mg/mL (3.61 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (3.61 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Maytansine is a highly potent microtubule-targeted compound that induces mitotic arrest and kills tumor cells at subnanomolar concentrations^[1].

In Vitro

Maytansine, at 6x10⁻⁸ M, irreversibly inhibits cell division in eggs of sea urchins and clams. Maytansine causes the disappearance of a mitotic apparatus or prevents one from forming if added at early stages. Maytansine inhibits in vitro polymerization of tubulin^[2].

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

REFERENCES

[1]. Lopus M, et al. Maytansine and cellular metabolites of antibody-maytansinoid conjugates strongly suppress microtubule dynamics by binding to microtubules. Mol Cancer Ther. 2010;9(10):2689-2699.

[2]. Remillard S, et al. Antimitotic activity of the potent tumor inhibitor maytansine. Science. 1975;189(4207):1002-1005.

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

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