Inhibitors

Demecolcine

Cat. No.: HY-N0282 CAS No.: 477-30-5 Molecular Formula: $\mathsf{C}_{21}\mathsf{H}_{25}\mathsf{NO}_5$ Molecular Weight: 371.43

Microtubule/Tubulin; Apoptosis Target:

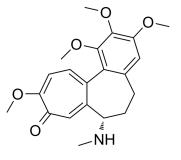
Pathway: Cell Cycle/DNA Damage; Cytoskeleton; Apoptosis

Storage: Powder -20°C 3 years

4°C 2 years

-80°C 6 months In solvent

-20°C 1 month



Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 25 mg/mL (67.31 mM; ultrasonic and warming and heat to 60°C) H₂O: 5 mg/mL (13.46 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.6923 mL	13.4615 mL	26.9230 mL
	5 mM	0.5385 mL	2.6923 mL	5.3846 mL
	10 mM	0.2692 mL	1.3461 mL	2.6923 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 (6.73 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.73 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.73 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Demecolcine is a potent mitotic inhibitor with an IC ₅₀ value of 2.4 μ M for inhibition of tubulin polymerization. Colcemid (Demecolcine) can interact with tubulin dimers to induce anti-mitotic action and inhibit microtubule growth. Colcemid (Demecolcine) can be used for inflammatory disorders and cancer research ^{[1][2]} .
In Vitro	Demecolcine (0.1-0.25 μ g/ml, 1 h) reduces the the hypoploidy frequencies in metaphase II complements of mouse, rat and frog spermatocytes ^[3] .

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

In Vivo

Demecolcine (0.3 mg/kg for intraperitoneal injection) increases the incidence of metaphase II hypoploidy in demecolcine treated mice $^{[3]}$.

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

Animal Model:	Mice ^[3]	
Dosage:	0.3 mg/kg	
Administration:	Intraperitoneal injection (i.p.)	
Result:	Elevated 7.8-8-fold of the hypoploid and hyperploid frequencies in metaphase II cells relative to controls.	

CUSTOMER VALIDATION

- Protein Cell. 2022 Jan;13(1):47-64.
- Cancer Lett. 2022 Aug 20;215855.
- Oncol Lett. 2023 Apr 19,25(6): 1-6.

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REFERENCES

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- [2]. Muzaffar A, et al. Antitubulin effects of derivatives of 3-demethylthiocolchicine, methylthio ethers of natural colchicinoids, and thioketones derived from thiocolchicine. Comparison with colchicinoids. J Med Chem. 1990 Feb;33(2):567-71.
- [3]. Risley MS, et al. An improved method for cytogenetic analysis of meiotic aneuploidy in rodent and frog spermatocytes. Mutat Res. 1990 Dec;234(6):361-8.
- [4]. T Tsuchida, et al. Colcemid-induced apoptosis of cultured human glioma: electron microscopic and confocal laser microscopic observation of cells sorted in different phases of cell cycle. Cytometry. 1998 Apr 1;31(4):295-9.
- [5]. Ashley M Rozario, et al. Ultra-Low Colcemid Doses Induce Microtubule Dysfunction as Revealed by Super-Resolution Microscopy. Bioexiv.

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

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