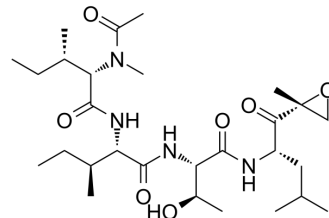


Epoxomicin

Cat. No.:	HY-13821
CAS No.:	134381-21-8
Molecular Formula:	C ₂₈ H ₅₀ N ₄ O ₇
Molecular Weight:	554.72
Target:	Proteasome; Apoptosis
Pathway:	Metabolic Enzyme/Protease; 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 (180.27 mM; Need ultrasonic)																					
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent</th> <th rowspan="2">Mass</th> <th colspan="3">Concentration</th> </tr> <tr> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> </thead> <tbody> <tr> <td rowspan="3">Preparing Stock Solutions</td> <td>1 mM</td> <td>1.8027 mL</td> <td>9.0136 mL</td> <td>18.0271 mL</td> </tr> <tr> <td>5 mM</td> <td>0.3605 mL</td> <td>1.8027 mL</td> <td>3.6054 mL</td> </tr> <tr> <td>10 mM</td> <td>0.1803 mL</td> <td>0.9014 mL</td> <td>1.8027 mL</td> </tr> </tbody> </table>	Solvent	Mass	Concentration			1 mg	5 mg	10 mg	Preparing Stock Solutions	1 mM	1.8027 mL	9.0136 mL	18.0271 mL	5 mM	0.3605 mL	1.8027 mL	3.6054 mL	10 mM	0.1803 mL	0.9014 mL	1.8027 mL
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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 (4.51 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.51 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.51 mM); Clear solution 																					

BIOLOGICAL ACTIVITY

Description	Epoxomicin (BU-4061T) is an epoxyketone-containing natural product and a potent, selective and irreversible proteasome inhibitor. Epoxomicin covalently binds to the LMP7, X, MECL1, and Z catalytic subunits of the proteasome and potently inhibits primarily the chymotrypsin-like activity. Epoxomicin can cross the blood-brain barrier. Epoxomicin has strongly antitumor and anti-inflammatory activity ^{[1][2][3][4][5]} .
IC₅₀ & Target	Proteasome ^[1]

In Vitro	<p>Epoxomicin shows quite potent cytotoxicities against all of the cells tested. Epoxomicin inhibits the cells growth of B16-F10, HCT116, Moser, P388 and K562 cells of IC₅₀ values of 0.002 µg/mL, 0.005 µg/mL, 0.044 µg/mL, 0.002 µg/mL and 0.037 µg/mL [1].</p> <p>Epoxomicin has antiproliferative activity with an IC₅₀ of 4 nM in EL4 lymphoma cells[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>								
In Vivo	<p>Epoxomicin (0.063-1 mg/kg; intraperitoneal injection; once daily; for 9 days; male BDFX mice) treatment shows significant antitumor effect with the minimum effective dose of 0.13mg/kg/day^[1].</p> <p>Epoxomicin also effectively inhibits NF-κB activation in vitro and potently blocks in vivo inflammation in the murine ear edema assay^[3].</p> <p>Epoxomicin is injected into adult rats over a period of 2 weeks. After a latency of 1 to 2 weeks, animals developed progressive Parkinsonism with bradykinesia, rigidity, tremor, and an abnormal posture. Postmortem analyses shows striatal dopamine depletion and dopaminergic cell death with apoptosis in the substantia nigra pars compacta^[4].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1" data-bbox="347 625 1515 863"> <tr> <td data-bbox="347 625 618 688">Animal Model:</td> <td data-bbox="618 625 1515 688">Male BDFX mice with B16 melanoma^[1]</td> </tr> <tr> <td data-bbox="347 688 618 751">Dosage:</td> <td data-bbox="618 688 1515 751">0.063 mg/kg, 0.13 mg/kg, 0.25 mg/kg, 0.5 mg/kg, 1 mg/kg</td> </tr> <tr> <td data-bbox="347 751 618 814">Administration:</td> <td data-bbox="618 751 1515 814">Intraperitoneal injection; once daily; for 9 days</td> </tr> <tr> <td data-bbox="347 814 618 863">Result:</td> <td data-bbox="618 814 1515 863">Exhibited strong therapeutic activity against B16 melanoma.</td> </tr> </table>	Animal Model:	Male BDFX mice with B16 melanoma ^[1]	Dosage:	0.063 mg/kg, 0.13 mg/kg, 0.25 mg/kg, 0.5 mg/kg, 1 mg/kg	Administration:	Intraperitoneal injection; once daily; for 9 days	Result:	Exhibited strong therapeutic activity against B16 melanoma.
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CUSTOMER VALIDATION

- Nat Metab. 2022 Sep;4(9):1202-1213.
- Nat Commun. 2021 Feb 19;12(1):1172.
- Redox Biol. 2023 Apr 20;62:102706.
- Mol Plant Pathol. 2018 Dec;19(12):2623-2634.
- Norwegian University of Science and Technology. Department of Biotechnology and Food Science. 2021 Oct.

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- [1]. Kim KB, et al. Proteasome inhibition by the natural products epoxomicin and dihydroeponemycin: insights into specificity and potency. *Bioorg Med Chem Lett*. 1999 Dec 6;9(23):3335-40.
- [2]. Hanada M, et al. Epoxomicin, a new antitumor agent of microbial origin. *J Antibiot (Tokyo)*. 1992 Nov;45(11):1746-52.
- [3]. Garrett IR, et al. Selective inhibitors of the osteoblast proteasome stimulate bone formation in vivo and in vitro. *J Clin Invest*. 2003 Jun;111(11):1771-82.
- [4]. McNaught KS, et al. Systemic exposure to proteasome inhibitors causes a progressive model of Parkinson's disease. *Ann Neurol*. 2004 Jul;56(1):149-62.
- [5]. Meng L, et al. Epoxomicin, a potent and selective proteasome inhibitor, exhibits in vivo antiinflammatory activity. *Proc Natl Acad Sci U S A*. 1999 Aug 31;96(18):10403-8.

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

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