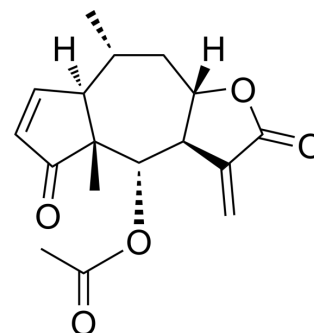


Bigelovin

Cat. No.:	HY-116506
CAS No.:	3668-14-2
Molecular Formula:	C ₁₇ H ₂₀ O ₅
Molecular Weight:	304.34
Target:	RAR/RXR; Apoptosis; Reactive Oxygen Species; Autophagy
Pathway:	Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor; Apoptosis; Immunology/Inflammation; NF-κB; Autophagy
Storage:	4°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (328.58 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	3.2858 mL	16.4290 mL	32.8580 mL
		5 mM	0.6572 mL	3.2858 mL	6.5716 mL
		10 mM	0.3286 mL	1.6429 mL	3.2858 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 (8.21 mM); Clear solution				
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (8.21 mM); Clear solution				
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (8.21 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Bigelovin, a sesquiterpene lactone isolated from <i>Inula hupehensis</i> , is a selective retinoid X receptor α agonist. Bigelovin suppresses tumor growth through inducing apoptosis and autophagy via the inhibition of mTOR pathway regulated by ROS generation ^[1] .
In Vitro	Bigelovin (0-20 μM, 24-72 h) significantly inhibits cell viability of liver cancer cells and induces apoptosis and autophagy ^[1] . Bigelovin causes a significant increase of p62, LC3B-II, Beclin-1 and a corresponding decrease of p62 levels in a time-dependent manner ^[1] . Bigelovin induces cell death involves the suppression of mTOR pathway regulated by ROS production ^[1] .

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

Cell Viability Assay^[1]

Cell Line:	HepG2 and SMMC-7721 cells.
Concentration:	0-20 μ M.
Incubation Time:	24, 48, 72 h.
Result:	Significantly reduced the cell viability of HepG2 and SMMC-7721 cells in a dose- and time-dependent manner. No significant difference observed in cell viability of normal liver cell lines, LO2 and LX2, after BigV treatment for 24, 48 or 72 h.

Western Blot Analysis^[1]

Cell Line:	HepG2 and SMMC-7721 cells.
Concentration:	0-10 μ M.
Incubation Time:	24 h.
Result:	The expression of Bcl-2 was decreased, whereas Bax was increased after treatment with BigV. Moreover, Caspase-9, -3 and PARP cleavage were activated significantly after BigV treatment.

In Vivo

Bigelovin (BigV, 5, 10, 20 mg/kg) exerts anti-tumor activity in HepG2 xenograft tumor model^[1].

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

Animal Model:	HepG2 xenograft model based on the male athymic BALB/c nude mice (5-6 weeks old, 18-22 g) ^[1] .
Dosage:	5, 10, 20 mg/kg.
Administration:	Intravenous injection every 2 days.
Result:	The tumor growth rate was significantly slower in BigV treated groups in a dose-dependent manner, along with the reduced tumor weight. No significant alteration of body weight and hepatic enzyme levels (AST, ALT and LDH) in serum was observed after BigV administration. Western blot findings of tumor tissues indicated the activation of apoptosis and autophagy characterized by the increase of cleaved Caspase-3 and PARP, as well as LC3BII levels. The inactivation of mTOR was also observed in tumor tissues isolated from BigV-treated mice.

REFERENCES

[1]. Bei Wang, et al. Bigelovin, a sesquiterpene lactone, suppresses tumor growth through inducing apoptosis and autophagy via the inhibition of mTOR pathway regulated by ROS generation in liver cancer. *Biochem Biophys Res Commun.* 2018 May 5;499(2):156-163.

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

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