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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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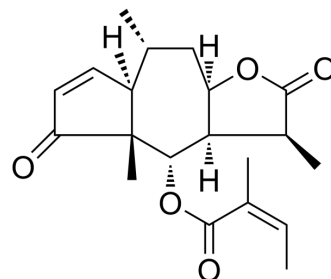
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Brevilin A

Cat. No.:	HY-N2959		
CAS No.:	16503-32-5		
Molecular Formula:	C ₂₀ H ₂₆ O ₅		
Molecular Weight:	346.42		
Target:	JAK; STAT; Apoptosis; Autophagy		
Pathway:	Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt; Apoptosis; Autophagy		
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 (288.67 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
	Preparing Stock Solutions	1 mM	2.8867 mL	14.4333 mL
	5 mM	0.5773 mL	2.8867 mL	5.7733 mL
	10 mM	0.2887 mL	1.4433 mL	2.8867 mL
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 >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 10 mg/mL (28.87 mM); Clear solution Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 6.25 mg/mL (18.04 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 6.25 mg/mL (18.04 mM); Clear solution 			

BIOLOGICAL ACTIVITY

Description	Brevilin A is an orally active STAT3/JAK inhibitor (STAT3 IC ₅₀ =?10.6 μM). Brevilin A shows anti-tumor activity, anti-proliferative activity to cancer cells, and can induce apoptosis and autophagy ^{[1][2]} .
IC₅₀ & Target	STAT3 10.6 μM (IC ₅₀)

In Vitro

Brevilin A (1-20 μM ; 24 h) inhibits STAT3 signaling in a dose dependent manner^[1].
Brevilin A (0-50 μM ; 24-72 h) inhibits the proliferation of NPC cells^[2].
Brevilin A (10 μM ; 24 and 48 h) induces DU145 and MDA-MB-468 apoptosis after 24 h treatment^[1].
Brevilin A (12.5 and 25 μM ; 24 h) blocks STAT3 tyrosine 705 phosphorylation in A549R cells^[1].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.
Cell Viability Assay^[1]

Cell Line:	A549R cells
Concentration:	1-20 μM
Incubation Time:	24 hours
Result:	Exhibited STAT3 signaling inhibition in a dose dependent manner with IC ₅₀ value of 10.6 μM .

Cell Proliferation Assay^[2]

Cell Line:	CNE-2 cells
Concentration:	0-50 μM
Incubation Time:	24, 48, and 72 hours
Result:	Showed IC ₅₀ values in CNE-2 cells with treatment times of 24, 48, and 72 h of 7.93, 2.60, and 22.26 μM , respectively.

Apoptosis Analysis^[1]

Cell Line:	DU145 and MDA-MB-468 cells
Concentration:	10 μM
Incubation Time:	24 and 48 hours
Result:	Decreased c-Myc and CyclinD1 after 24 h and 48 h treatment, increased cleaved PARP after 24 h treatment.

Western Blot Analysis^[1]

Cell Line:	A549R cells
Concentration:	12.5 and 25 μM
Incubation Time:	24 hours
Result:	Inhibits STAT3 phosphorylation in A549R cells.

In Vivo

Brevilin A (oral gavage; 10 and 20 mg/kg; once daily; 16 d) inhibits CNE-2 xenograft tumor growth, and inhibits PI3K/AKT and STAT3 signaling in vivo^[2].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male BALB/c nude mice injected with CNE-2 cells ^[2]
Dosage:	10 and 20 mg/kg
Administration:	Oral gavage; 10 and 20 mg/kg; once daily; 16 days

Result:

Decreased average tumor volumes and weights treated with 20 mg/kg by 36.3% and 46.0%, respectively, compared to vehicle control.

Inhibited the protein expression of p-AKT and p-STAT3 at both low and high doses.

CUSTOMER VALIDATION

- Cell Cycle. 2024 Jan 25:1-13.
- Mol Biotechnol. 2024 May 14.
- Ann Med Surg (Lond). 2023 Oct 18.
- Research Square Preprint. 2024 Mar 6.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Chen X, et al. Brevilin A, a novel natural product, inhibits janus kinase activity and blocks STAT3 signaling in cancer cells. PLoS One. 2013 May 21;8(5):e63697.

[2]. You P, et al. Brevilin A induces apoptosis and autophagy of colon adenocarcinoma cell CT26 via mitochondrial pathway and PI3K/AKT/mTOR inactivation. Biomed Pharmacother. 2018 Feb;98:619-625.

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

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