

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

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- Trockeneiszuschlag
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SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

linkedin.com/company/szaboscandic in



Proteins

Product Data Sheet

Vanicoside B

Cat. No.: HY-N9561 CAS No.: 155179-21-8 Molecular Formula: $C_{49}H_{48}O_{20}$ Molecular Weight: 956.89 Target: CDK; STAT

Pathway: Cell Cycle/DNA Damage; JAK/STAT Signaling; Stem Cell/Wnt

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 (104.51 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.0451 mL	5.2253 mL	10.4505 mL
	5 mM	0.2090 mL	1.0451 mL	2.0901 mL
	10 mM	0.1045 mL	0.5225 mL	1.0451 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 (2.61 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (2.61 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (2.61 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Vanicoside B is a phenylpropanoyl sucrose derivative, can be isolated from the herb Persicaria dissitiflora. Vanicoside B targets cyclin-dependent kinase 8 (CDK8) and exhibits anti-tumor activity. The potential mechanism is Vanicoside B blocks CDK8-mediated signaling pathways and decreases the expression of epithelial-mesenchymal transition proteins, so that it leads to cell cycle arrest and apoptosis^{[1][2]}.

IC₅₀ & Target

CDK3

STAT3

In Vitro

Vanicoside B (2.5-20 μM; 72 h) shows antiproliferative activity against a panel of cancer cell lines in triple-negative breast cancer (TNBC) MDA-MB-231 cells and HCC38 cells^[1].

Vanicoside B (2.5-20 μ M; 72 h, 14 d, and 72 h, respectively) inhibits cell viability, colony formation, and disturbs cell cycle distribution in TNBC cells^[1].

Vanicoside B (2.5-10 μ M; 48 h) decreased p-STAT1, p-STAT3, and p-S6 protein level, and induces apoptosis by regulating the Skp2-p27 axis in TNBC cells^[1].

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

Western Blot Analysis^[1]

Incubation Time:

Result:

Cell Line:	MDA-MB-231 cells and HCC38 cells	
Concentration:	0, 2.5, 5, 10 μΜ	
Incubation Time:	48 hours	
Result:	Increased cleaved PARP, and p27 protein expressions, but decreased Skp2 protein level. Suppressed CDK8 target genes and the expression of EMT-associated proteins. Suppressed the expression of the cell proliferation marker Ki-67 in tumor tissues, also significantly suppressed the expressions of p-STAT1 (S727) and AXL.	
Cell Cycle Analysis ^[1]		
Cell Line:	MDA-MB-231 cells and HCC38 cells	
Concentration:	0, 2.5, 5, 10 μΜ	

In Vivo

Vanicoside B (5 mg/kg and 20 mg/kg; i.p.; 3 times per week for 4 weeks) inhibits tumor growth in xenografted mouse models with MDAMB-231 cells^[1].

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

Inhibited cell cycle at sub-G1 phase.

72 hours

Animal Model:	MDA-MB-231 cell-implanted xenograft mouse model $^{[1]}$
Dosage:	5 mg/kg, 20 mg/kg
Administration:	Intraperitoneal injection; 3 times per week over 4 weeks
Result:	Significantly reduced tumor volumes at 5 mg/kg and 20 mg/kg by 53.85% and 65.72%, respectively.

REFERENCES

[1]. Kim D, et al. Antitumor Activity of Vanicoside B Isolated from Persicaria dissitiflora by Targeting CDK8 in Triple-Negative Breast Cancer Cells. J Nat Prod. 2019 Nov 22;82(11):3140-3149.

[2]. Takasaki M, et al. Cancer chemopreventive activity of phenylpropanoid esters of sucrose, vanicoside B and lapathoside A, from Polygonum lapathifolium. Cancer Lett. 2001 Nov 28;173(2):133-8.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA