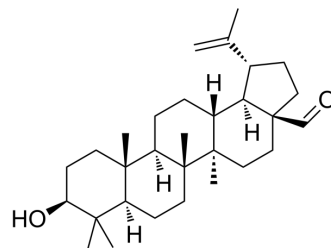


## Betulinaldehyde

<b>Cat. No.:</b>	HY-N0084		
<b>CAS No.:</b>	13159-28-9		
<b>Molecular Formula:</b>	C <sub>30</sub> H <sub>48</sub> O <sub>2</sub>		
<b>Molecular Weight:</b>	440.7		
<b>Target:</b>	Bacterial		
<b>Pathway:</b>	Anti-infection		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 20 mg/mL (45.38 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	2.2691 mL	11.3456 mL	22.6912 mL
		5 mM	0.4538 mL	2.2691 mL	4.5382 mL
10 mM		0.2269 mL	1.1346 mL	2.2691 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2 mg/mL (4.54 mM); Clear solution  2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2 mg/mL (4.54 mM); Clear solution				

### BIOLOGICAL ACTIVITY

<b>Description</b>	<p>Betulinaldehyde (Betunal) Has anti-cancer and anti-staphylococcus aureus activity. Betulinaldehyde Suppressible Akt, MAPK sum STAT3 Signal path, increase self-transfer, Suppression A549 Cellular vitality, increase and transfer. Betulinaldehyde suppresses PLCγ1/Ca<sup>2+</sup>/MMP9 signal pathway, has a key effect on vascular plasticity, and is available for cardiovascular disease (CVD) research.</p>
<b>In Vitro</b>	<p>Betulinaldehyde (20, 40, 80 μM; 24 h) inhibits the colony formation ability of A549 cells in a concentration-dependent manner<sup>[3]</sup>.</p> <p>Betulinaldehyde (20 μM; 24 h) inhibits the phosphorylation of Akt, MAPK and STAT3 in A549 cells<sup>[3]</sup>.</p> <p>Betulinaldehyde (80 μM, 160 μM; 24 h) leads vascular remodeling in rats with carotid balloon injurys, and inhibits intracellular Ca<sup>2+</sup> levels<sup>[4]</sup>.</p>

---

	MCE has not independently confirmed the accuracy of these methods. They are for reference only.
<b>In Vivo</b>	Betulinoldehyde (50-200 mg/kg; ip; once daily for 1 week) inhibits the growth of A549 tumor xenografts in nude mice in vivo <sup>[3]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

---

## CUSTOMER VALIDATION

- Phytomedicine. 2023 May 18, 154891.
- Fuel. 2018 Dec 15, 234:110-119.
- Catalysis Today. 2020 Aug.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

## REFERENCES

- [1]. Chung PY, et al. Identification, by gene expression profiling analysis, of novel gene targets in *Staphylococcus aureus* treated with betulinaldehyde. *Res Microbiol*. 2013 May;164(4):319-26.
- [2]. Chung PY, et al. Transcriptional profiles of the response of methicillin-resistant *Staphylococcus aureus* to pentacyclic triterpenoids. *PLoS One*. 2013;8(2):e56687.
- 

**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](mailto:tech@MedChemExpress.com)

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