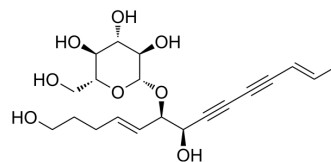


Lobetyolin

Cat. No.:	HY-N0327
CAS No.:	129277-38-9
Molecular Formula:	C ₂₀ H ₂₈ O ₈
Molecular Weight:	396.43
Target:	Apoptosis; Xanthine Oxidase
Pathway:	Apoptosis; Metabolic Enzyme/Protease
Storage:	-20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (126.13 mM; Need ultrasonic)
H₂O : 50 mg/mL (126.13 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	2.5225 mL	12.6126 mL	25.2251 mL
	5 mM	0.5045 mL	2.5225 mL	5.0450 mL
	10 mM	0.2523 mL	1.2613 mL	2.5225 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: PBS
Solubility: 33.33 mg/mL (84.08 mM); Clear solution; Need ultrasonic
- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Lobetyolin, a bioactive compound, is derived from *Codonopsis pilosula*. Lobetyolin has anti-inflammatory, anti-oxidative and xanthine oxidase inhibiting activities. Lobetyolin also induces the apoptosis via the inhibition of ASCT2-mediated glutamine metabolism^{[1][2]}. Lobetyolin is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAC) with molecules containing Azide groups.

In Vitro

Lobetyolin is derived from *Codonopsis pilosula* and has antioxidative effect. MUC5AC gene expression induced by phorbol

12-myristate 13-acetate (PMA) from NCI-H292 cells is inhibited by pretreatment with Lobetyolin, lobetyol or methyl linoleate. Lobetyolin does not affect PMA-induced MUC5AC production from NCI-H292 cells. The amounts of mucin in the cells of Lobetyolin-treated cultures are 100±25%, 487±33%, 524±38%, 411±24%, and 402±45% for control, 10 ng/mL of PMA alone, PMA plus Lobetyolin 1 µM, PMA plus Lobetyolin 10 µM, and PMA plus Lobetyolin 100 µM, respectively^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Cell Assay ^[1]

After 24 hours of serum deprivation, cells are pretreated with Lobetyolin, lobetyol or methyl linoleate (1, 10, and 100 µM), for 30 minutes and then treated with phorbol 12-myristate 13-acetate (PMA; 10 ng/mL) for 24 hours in serum-free RPMI 1640. After 24 hours, the spent media are collected to measure the secretion of MUC5AC protein and cells are lysed with buffer solution containing 20 mM Tris, 0.5% NP-40, 250 mM NaCl, 3 mM EDTA, 3 mM EGTA, and protease inhibitor cocktail and collected to measure the production of MUC5AC protein (in 24-well culture plate)^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Front Pharmacol. 2024 May 10.

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REFERENCES

[1]. Yoon YP, et al. Effects of Lobetyolin, Lobetyol and Methyl linoleate on Secretion, Production and Gene Expression of MUC5AC Mucin from Airway Epithelial Cells. *Tuberc Respir Dis (Seoul)*. 2014 Nov;77(5):203-8.

[2]. He W, et, al. Lobetyolin induces apoptosis of colon cancer cells by inhibiting glutamine metabolism. *J Cell Mol Med*. 2020 Mar;24(6):3359-3369.

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

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