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- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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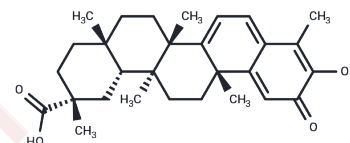
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Tripterin

Chemical Properties

CAS No. :	34157-83-0
Formula:	C ₂₉ H ₃₈ O ₄
Molecular Weight:	450.61
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	Tripterin (Celastrol) is a natural product, a proteasome inhibitor that inhibits the pancreatic rennet-like activity of the 20S proteasome (IC ₅₀ =2.5 μM). Tripterin has anti-inflammatory, anti-infectious, and immunomodulatory properties.
Targets(IC ₅₀)	Apoptosis, Mitophagy, Proteasome, Autophagy
In vitro	<p>METHODS: Human prostate cancer cells PC-3 were treated with Tripterin (0.5-5 μM) for 12 h. Proteasomal chymotrypsin-like activity was assayed using Z-GGL-AMC.</p> <p>RESULTS: Tripterin significantly inhibited proteasomal chymotrypsin-like activity in PC-3 cells in a concentration-dependent manner, reaching about 55% inhibition at 2.5 μM. [1]</p> <p>METHODS: Human chronic myeloid leukemia cells KBM-5 were incubated with Tripterin (2.5 μM) for 6 h, followed by treatment with TNF (1 nM) for 6-24 h. Target protein expression levels were detected using Western Blot.</p> <p>RESULTS: TNF induced the expression of anti-apoptotic proteins IAP1, IAP2, Bcl-2, Bcl-XL, c-FLIP and survivin in a time-dependent manner, which was inhibited by Tripterin. [2]</p>
In vivo	<p>METHODS: To detect anti-tumor activity in vivo, Tripterin (1-3 mg/kg, 10% DMSO+70% Cremophor/ethanol (3:1)+20% PBS) was injected intraperitoneally once daily for sixteen days into nude immunodeficient mice bearing human prostate cancer tumor PC-3.</p> <p>RESULTS: Tripterin treatment significantly inhibited the growth of prostate cancer xenografts and suppressed proteasome activity and induced apoptosis in vivo. [1]</p> <p>METHODS: To detect anti-tumor activity in vivo, Tripterin (1.25 mg/kg) was intraperitoneally injected into BALB/c (nu/nu) mice bearing vestibular nerve sheath tumor SC4 every three days for two weeks.</p> <p>RESULTS: Tripterin significantly inhibited tumor growth without showing toxicity. [3]</p>
Kinase Assay	Inhibition of purified 20S proteasome activity: A purified rabbit 20S proteasome (0.1 μg) is incubated with 40 μM of various fluorogenic peptide substrates in 100 μL assay buffer (20 mM Tris-HCl (pH 7.5)), in the presence of Celastrol at different concentrations or in the solvent DMSO for 2 hours at 37 °C, followed by measurement of inhibition of each proteasomal activity.
Cell Research	The anti-proliferative effect of celastrol on various human tumor cell lines is determined by the MTT uptake method. Briefly, 5×10 ³ cells are incubated with Celastrol in triplicate in a 96-well plate at 37 °C. MTT solution is then added to each well. After a 2 hours incubation at 37 °C, extraction buffer (20% SDS, 50% dimethylformamide) is added, cells are incubated overnight at 37 °C, and the optical density is then measured at 570 nm using a Tecan plate reader.(Only for Reference)

Solubility Information

Solubility	DMSO: 60 mg/mL (133.15 mM), Ethanol: 33.8 mg/mL (75 mM), (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.2192 mL	11.0961 mL	22.1921 mL
5 mM	0.4438 mL	2.2192 mL	4.4384 mL
10 mM	0.2219 mL	1.1096 mL	2.2192 mL
50 mM	0.0444 mL	0.2219 mL	0.4438 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Reference

Yang H, et al. Celastrol, a triterpene extracted from the Chinese "Thunder of God Vine," is a potent proteasome inhibitor and suppresses human prostate cancer growth in nude mice. *Cancer Res.* 2006 May 1;66(9):4758-65.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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