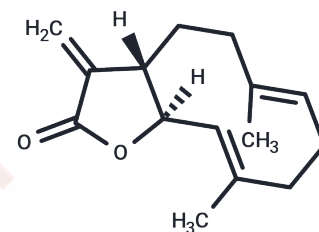


Costunolide

Chemical Properties

CAS No. :	553-21-9
Formula:	C ₁₅ H ₂₀ O ₂
Molecular Weight:	232.32
Storage:	Store at low temperature Powder: -20°C for 3 years In solvent: -80°C for 1 year <small>Actual storage temperature shall be subject to the COA.</small>



Biological Description

Description	Costunolide (Costus lactone) has anti-inflammatory and anti-oxidant properties and mediates apoptosis.
Targets(IC50)	Apoptosis,Endogenous Metabolite,Telomerase
In vitro	Costunolide reduces neovascularization induced by ethyl ester resin scale plaster in the corneal micropocket model of mice. It inhibits angiogenesis by blocking the vascular endothelial growth factor signaling pathway.
In vivo	Costunolide has been shown to inhibit cell proliferation in a dose-dependent manner over a 48-hour treatment in various human tumor cells, including A549, SK-OV-3, SK-MEL-2, XF498, and HCT-15. It induces apoptosis in HL-60 human leukemia cells through ROS-mediated mitochondrial permeability transition and cytochrome C release. Additionally, Costunolide exhibits significant antifungal activity against organisms such as Trichophyton rubrum, T. mentagrophytes, and Microsporum gypseum. The compound also suppresses telomerase activity and cell proliferation in MCF-7 and MDA-MB-231 cells in a concentration and time-dependent manner. Furthermore, Costunolide dose-dependently inhibits the farnesylation of lamin B by farnesyl protein transferase, demonstrating its diverse biochemical impacts.
Kinase Assay	Telomerase activity assay: The telomerase activity is measured by the TRAP assay using the TRAPez Telomerase Detection Kit, which includes primers of a 36-bp internal control (IC) for quantifying the amplification of telomerase activity within a linear range close to 2.5 logs. For RNase treatment, 10µL of extract are incubated with 1µg of RNase at 37 °C for 20 minutes. The products of the TRAP assay are resolved by electrophoresis in a non-denaturing 12% PAGE in a buffer containing 0.5 × Tris-borate EDTA and detected by autoradiograph. For quantification of TRAP products, the dried gels are exposed to Fuji Imaging Plate at room temperature. Results are corrected for background, and a standard value of 100 is given to the untreated control cell signal. Signal intensities of Costunolide-treated cells are compared to the standard and are expressed as a fraction of the maximum value of 100. [1]
Cell Research	1) Plate 500-10,000 cells in 200 µL media per well in a 96 well plate. Leave 8 wells empty for blank controls. 2) Incubate (37 °C, 5% CO ₂) overnight to allow the cells to attach to the wells. 3) Add 2 µL of Costunolide dissolved in DMSO to each well. Place on a shaking table, 150 rpm for 5 minutes, to thoroughly mix the samples into the media. 5) Incubate

Cell Research	(37 °C, 5% CO ₂) for 48 hours to allow Costunolide to take effect. 6) Make 2 mL or more of MTT solution per 96 well plate at 5 mg/mL in PBS. Do not make a stock as MTT in solution is not stable long-term. 7) Add 20 µL MTT solution to each well. Place on a shaking table, 150 rpm for 5 minutes, to thoroughly mix the MTT into the media. 8) Incubate (37 °C, 5% CO ₂) for 1-5 hours to allow the MTT to be metabolized. 9) Dump off the media. (Dry plate on paper towels to remove residue if necessary. 10) Resuspend formazan (MTT metabolic product) in 200 µL DMSO. Place on a shaking table, 150 rpm for 5 minutes, to thoroughly mix the formazan into the solvent. 11) Read optical density at 560 nm and subtract background at 670 nm. Optical density should be directly correlated with cell quantity. (Only for Reference)
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Solubility Information

Solubility	DMSO: 240 mg/mL (1033.06 mM),Sonication is recommended. Ethanol: 11.6 mg/mL (49.93 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+40% PEG300+5% Tween 80+45% Saline: 5 mg/mL (21.52 mM),Sonication is recommended. <i>Please add the solvents sequentially, clarifying the solution as much as possible before adding the next one. Dissolve by heating and/or sonication if necessary. Working solution is recommended to be prepared and used immediately. The formulation provided above is for reference purposes only. In vivo formulations may vary and should be modified based on specific experimental conditions.</i>

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	4.3044 mL	21.522 mL	43.0441 mL
5 mM	0.8609 mL	4.3044 mL	8.6088 mL
10 mM	0.4304 mL	2.1522 mL	4.3044 mL
50 mM	0.0861 mL	0.4304 mL	0.8609 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.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

- Choi SH, et al, Cancer Lett, 2005, 227(2), 153-162.
Park SH, et al, Planta Med, 2001, 67(4), 358-359.
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Jeong SJ, et al, Cancer Lett, 2002, 187(1-2), 129-133.

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