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Zuschläge

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- Expressversand

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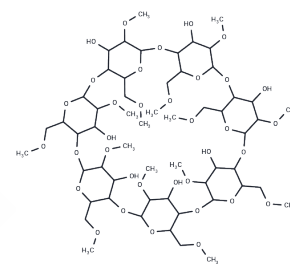
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Methyl- β -cyclodextrin

Chemical Properties

CAS No. :	128446-36-6
Formula:	C54H94O35
Molecular Weight:	
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	Methyl- β -cyclodextrin (Methyl-beta-cyclodextrin) is a macrocyclic compound used as a solubilizer for hydrophobic compounds in biological experiments. Methyl- β -cyclodextrin is also a lipid raft inhibitor with cholesterol-lowering activity and potential antitumor activity.
Targets(IC50)	Others
In vitro	<p>METHODS: Five primary exudative lymphoma (PEL) cell lines, BCBL-1, BC-1, BC-3, TY-1, and GTO, were treated with Methyl-β-cyclodextrin (0-10 mM) for 24 h, and cell viability was measured by MTT assay.</p> <p>RESULTS: Methyl-β-cyclodextrin dose-dependently inhibited the growth of PEL cells with IC50 between 3.33-4.23 mM. [1]</p> <p>METHODS: Chicken hepatocellular carcinoma cells, LMH, were treated with Methyl-β-cyclodextrin (10 mM) for 1 h, then cholesterol (50 μg/mL) was added at different time points before, during, and after infection with FAdV-4, and the expression levels of the target proteins were detected by Western Blot.</p> <p>RESULTS: When LMH cells were pretreated with increasing concentrations of Methyl-β-cyclodextrin prior to infection, penton protein levels decreased in a dose-dependent manner. [2]</p>
In vivo	<p>METHODS: To test the antitumor activity in vivo, Methyl-β-cyclodextrin (500 mg/kg) was intraperitoneally injected into NRJ mice bearing PEL tumor BCBL-1 once a day for twenty-one days.</p> <p>RESULTS: Methyl-β-cyclodextrin significantly inhibited the growth and invasion of PEL cells without significant adverse effects. [1]</p> <p>METHODS: To investigate the modulatory effects on collagen, Methyl-β-cyclodextrin (1.25-5.0 mg/mouse) was intradermally injected into the SKH1 mouse twice a week for two months.</p> <p>RESULTS: Methyl-β-cyclodextrin showed potent COL I up-regulatory activity resulting in increased skin thickness. [3]</p>
Kinase Assay	PEL cells are incubated in triplicate in a 96-well microculture plate in the presence of different concentrations of methyl- β -cyclodextrin (0-10 mM) in a final volume of 0.1 mL for 24 h at 37°C. Subsequently, MTT (0.5 mg/mL final concentration) is added to each well. After 3 h of additional incubation, 100 μ L of a 0.04 N HCl is added to dissolve the crystals. Absorption values at 570 nm are determined.

A DRUG SCREENING EXPERT

Cell Research	PEL cells are incubated in triplicate in a 96-well microculture plate in the presence of different concentrations of methyl- β -cyclodextrin (0-10 mM) in a final volume of 0.1 mL for 24 h at 37°C. Subsequently, MTT (0.5 mg/mL final concentration) is added to each well. After 3 h of additional incubation, 100 μ L of a 0.04 N HCl is added to dissolve the crystals. Absorption values at 570 nm are determined[1].
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Solubility Information

Solubility	DMSO: 55 mg/mL (41.98 mM), (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Reference

Lu J Y, Huang W T, Zhou K, et al. Microbial Lipopeptide Supramolecular Self-Assemblies as a Methuosis-Like Cell Death Inducer with In Vivo Antitumor Activity. *Small*. 2021: 2104034.
Gotoh K, et al. The antitumor effects of

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