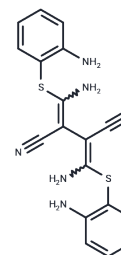


U0126

Chemical Properties

CAS No. : 109511-58-2
 Formula: C₁₈H₁₆N₆S₂
 Molecular Weight: 380.49
 Storage: Store at low temperature
 Powder: -20°C for 3 years | In solvent: -80°C for 1 year
 Actual storage temperature shall be subject to the COA.



Biological Description

Description	U0126, an effective and selective non-competitive inhibitor of MAP kinase, inhibits MEK-1 and MEK-2 with IC ₅₀ values of 0.07 and 0.06 μM respectively. U0126 inhibits autophagy and mitophagy.
Targets(IC ₅₀)	Mitophagy,MEK,Autophagy,Influenza Virus
In vitro	Rat hepatocarcinoma cells (FAO) stimulated by fetal calf serum (FCS) exhibits a significant proportion in S phase (32.62%) whereas U0126 strongly decreases the proportion of cells in S phase (9.92%) and increases the proportion of cells in G ₀ -G ₁ phase and to a lesser extent in G ₂ /M[1]. U0126 efficiently reduces progeny virus titers of all tested strains in A549 cells. While nM concentrations of U0126 are efficient to reduce H1N1v and H5N1 (MB1), μM concentrations of U0126 are required to reduce the virus titer of H5N1 (GSB) and H7N7. The EC ₅₀ values for U0126-EtOH against H1N1v are 1.2 μM in A549 cells and 74.7 μM in MDCKII cells[2].
In vivo	In mice, U0126-EtOH (U0126; i.p., 10.5 mg/kg) inhibits tumor growth with a 60-70% reduction 9 days after injection and thereafter[1]. After treatment with U0126-EtOH (U0126; i.p., 30 mg/kg), the vasoconstriction to S6c is markedly reduced in rats[3].

Solubility Information

Solubility	DMSO: 83.33 mg/mL (219.01 mM), H ₂ O: Soluble, (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.6282 mL	13.1409 mL	26.2819 mL
5 mM	0.5256 mL	2.6282 mL	5.2564 mL
10 mM	0.2628 mL	1.3141 mL	2.6282 mL
50 mM	0.0526 mL	0.2628 mL	0.5256 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

- Bessard A, et al. RNAi-mediated ERK2 knockdown inhibits growth of tumor cells in vitro and in vivo. *Oncogene*. 2008 Sep 11;27(40):5315-25.
- Li W Y, Shi T S, Huang J, et al. Activation of the mTORC1 signaling cascade in the hippocampus and medial prefrontal cortex is required for the antidepressant actions of vortioxetine in mice. *International Journal of Neuropsychopharmacology*. 2023: pyad017.
- Lu D, Zhang H, Zhang Y, et al. Secreted MbovP0145 Promotes IL-8 Expression through Its Interactive β -Actin and MAPK Activation and Contributes to Neutrophil Migration. *Pathogens* 2021, 10, 1628. *Pathogens*. 2021
- Xin J, Wang Z, Shen Y, et al. S100 calcium-binding protein A16 suppresses the osteogenic differentiation of rat bone marrow mesenchymal stem cells by inhibiting SMAD family member 4 signaling. *Experimental and Therapeutic Medicine*. 2024, 27(6): 1-10.
- Droebner K, et al. Antiviral activity of the MEK-inhibitor U0126 against pandemic H1N1v and highly pathogenic avian influenza virus in vitro and in vivo. *Antiviral Res.* 2011, 92(2), 195-203.
- Shen S, Huang Z, Lin L, et al. Tussilagone attenuates atherosclerosis through inhibiting MAPKs-mediated inflammation in macrophages. *International Immunopharmacology*. 2023, 119: 110066.
- Ahnstedt H, et al. U0126 attenuates cerebral vasoconstriction and improves long-term neurologic outcome after stroke in female rats. *J Cereb Blood Flow Metab.* 2015 Mar;35(3):454-60.
- Du P, Hu T, An Z, et al. In vitro and in vivo synergistic efficacy of ceritinib combined with programmed cell death ligand-1 inhibitor in anaplastic lymphoma kinase-rearranged non-small-cell lung cancer. *Cancer Science*. 2020, 111(6): 1887.
- Yang N, Zou C, Luo W, et al. Sclareol attenuates angiotensin II-induced cardiac remodeling and inflammation via inhibiting MAPK signaling. *Phytotherapy Research*. 2022
- Favata MF, et al. Identification of a novel inhibitor of mitogen-activated protein kinase kinase. *J Biol Chem*. 1998 Jul 17;273(29):18623-32.
- Du P, Hu T, An Z, et al. In vitro and in vivo synergistic efficacy of ceritinib combined with programmed cell death ligand-1 inhibitor in anaplastic lymphoma kinase-rearranged non-small-cell lung cancer[J]. *Cancer Science*. 2020, 111(6): 1887.
- Shi Y, Wang X, Meng Y, et al. A Novel Mechanism of Endoplasmic Reticulum Stress-and c-Myc-Degradation-Mediated Therapeutic Benefits of Antineurokinin-1 Receptor Drugs in Colorectal Cancer. *Advanced Science*. 2021: 2101936.
- Lu D, Zhang H, Zhang Y, et al. Secreted MbovP0145 Promotes IL-8 Expression through Its Interactive β -Actin and MAPK Activation and Contributes to Neutrophil Migration. *Pathogens*. 2021, 10(12): 1628.
- Shen S, Wu G, Luo W, et al. Leonurine attenuates angiotensin II-induced cardiac injury and dysfunction via inhibiting MAPK and NF- κ B pathway. *Phytomedicine*. 2022: 154519.
- Sun B, Wu H, Lu J, et al. Irisin reduces bone fracture by facilitating osteogenesis and antagonizing TGF- β /Smad signaling in a growing mouse model of osteogenesis imperfecta. *Journal of Orthopaedic Translation*. 2023, 38: 175-189.
- Wang D, Wang H, Yan Y, et al. Coating 3D-printed bioceramics with histatin promotes adhesion and osteogenesis of stem cells. *Tissue Engineering*. 2023 (ja).

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