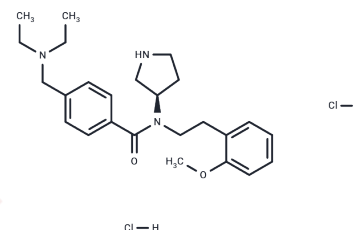


PF429242 dihydrochloride

Chemical Properties

CAS No. :	2248666-66-0
Formula:	C ₂₅ H ₃₇ Cl ₂ N ₃ O ₂
Molecular Weight:	482.49
Appearance:	no data available
Storage:	Powder: -20°C for 3 years In solvent: -80°C for 1 year



Biological Description

Description	PF429242 dihydrochloride can reversibly inhibit S1P (IC ₅₀ of 175 nM).
Targets(IC ₅₀)	S1P Receptor
In vitro	10 μM PF-429242 down-regulates the signal from an SRE-luciferase reporter gene in human embryonic kidney 293 cells and the expression of endogenous SREBP target genes in cultured HepG2 cells. In HepG2 cells, PF-429242 inhibits cholesterol synthesis, with an IC ₅₀ of 0.5 μM[1]. PF-429242 treatment also shows suppressive effects on DENV2 yields in the cultured fluids of human-derived HEK-293, Hep G2, and non-human-primate derived LLC-MK2 cells[2]. PF-429242 has potent antiviral against LCMV and LASV [3].
In vivo	PF-429242 inhibits the expression of hepatic SREBP target genes, and the hepatic rates of cholesterol and fatty acid synthesis are reduced In mice[1].

Solubility Information

Solubility	H ₂ O: 50 mg/mL (103.63 mM),Sonication is recommended. DMSO: 83.3 mg/mL (165.81 mM) (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.0726 mL	10.3629 mL	20.7258 mL
5 mM	0.4145 mL	2.0726 mL	4.1452 mL
10 mM	0.2073 mL	1.0363 mL	2.0726 mL
50 mM	0.0415 mL	0.2073 mL	0.4145 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

Hawkins JL, et al. Pharmacologic inhibition of site 1 protease activity inhibits sterol regulatory element-binding protein processing and reduces lipogenic enzyme gene expression and lipid synthesis in cultured cells and

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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