

Gue1654

Chemical Properties

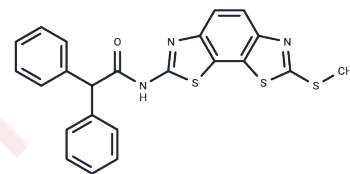
CAS No. : 397290-30-1

Formula: C₂₃H₁₇N₃O₃S₃

Molecular Weight: 447.6

Storage: 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	Gue1654 is a selective OXE-R inhibitor that attenuates coronary artery ligation-induced ischemic myocardial injury and cardiomyocyte oxygen/glucose deprivation-induced injury in mice through activation of BCAT1. Gue1654 inhibits OXE-R-inhibited protein kinase C-ε (PKC-ε)/nuclear factor κB (NF-κB) signaling and apoptosis in cardiomyocytes. Gue1654 can be used to study cardiovascular disease.
Targets(IC50)	Others,NF-κB,PKC

Solubility Information

Solubility	DMSO: 100 mg/mL (223.41 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
In vivo Formulation	10% DMSO+90% Corn Oil: 3.3 mg/mL (7.37 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	2.2341 mL	11.1707 mL	22.3414 mL
5 mM	0.4468 mL	2.2341 mL	4.4683 mL
10 mM	0.2234 mL	1.1171 mL	2.2341 mL
50 mM	0.0447 mL	0.2234 mL	0.4468 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

Blättermann S, et al. A biased ligand for OXE-R uncouples G α and G $\beta\gamma$ signaling within a heterotrimer. Nat Chem Biol. 2012;8(7):631-638.

Lai Q, et al. Oxoeicosanoid receptor inhibition alleviates acute myocardial infarction through activation of BCAT1. Basic Res Cardiol. 2021 Jan 23;116(1):3.

Cossette C, et al. Biosynthesis and actions of 5-oxoeicosatetraenoic acid (5-oxo-EET) on feline granulocytes. Biochem Pharmacol. 2015 Aug 1;96(3):247-55.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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