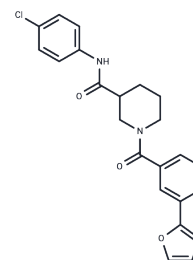


CCG-203971

Chemical Properties

CAS No. : 1443437-74-8
 Formula: C₂₃H₂₁ClN₂O₃
 Molecular Weight: 408.88
 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	CCG-203971 is an inhibitor of SRE activation in the prostate cancer cell line PC-3 (IC ₅₀ : 6.4 μM), with 87% inhibition of SRE activation achieved at 100 μM. This compound also inhibits PC-3 cell migration (IC ₅₀ : 4.2 μM), as determined by a scratch wound assay. CCG-203971 (CCG203971) causes no cytotoxicity when evaluated by the WST-1 assay. It is well tolerated in normal mice up to doses of 100 mg/kg given intraperitoneally over five days.
Targets(IC ₅₀)	Rho,Ras

Solubility Information

Solubility	DMSO: 50 mg/mL (122.29 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: 2 mg/mL (4.89 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	2.4457 mL	12.2285 mL	24.4571 mL
5 mM	0.4891 mL	2.4457 mL	4.8914 mL
10 mM	0.2446 mL	1.2229 mL	2.4457 mL
50 mM	0.0489 mL	0.2446 mL	0.4891 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

Haak AJ, et al. Targeting the myofibroblast genetic switch: inhibitors of myocardin-related transcription factor/serum response factor-regulated gene transcription prevent fibrosis in a murine model of skin injury. *J Pharmacol Exp Ther.* 2014 Jun;349(3):480-6.

Johnson LA, et al. Novel Rho/MRTF/SRF inhibitors block matrix-stiffness and TGF- β -induced fibrogenesis in human colonic myofibroblasts. *Inflamm Bowel Dis.* 2014 Jan;20(1):154-65.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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