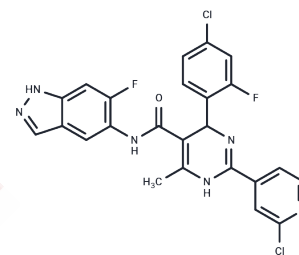


GSK-25

## Chemical Properties

CAS No. : 874119-56-9  
 Formula: C<sub>24</sub>H<sub>16</sub>Cl<sub>2</sub>F<sub>2</sub>N<sub>6</sub>O  
 Molecular Weight: 513.33  
 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	GSK-25 maintains good selectivity against a panel of 31 kinases, as well as RSK1 and p70S6K (RSK1 IC <sub>50</sub> of 398 nM, p70S6K IC <sub>50</sub> of 1000nM), and a dramatically improved P450 profile (>2.2 uM at all isozymes tested).
Targets(IC <sub>50</sub> )	mTOR,ROCK,S6 Kinase

## Solubility Information

Solubility	DMSO: 60 mg/mL (116.88 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 (3.9 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	1.9481 mL	9.7403 mL	19.4806 mL
5 mM	0.3896 mL	1.9481 mL	3.8961 mL
10 mM	0.1948 mL	0.974 mL	1.9481 mL
50 mM	0.039 mL	0.1948 mL	0.3896 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

Sehon CA, et al. Potent, selective and orally bioavailable dihydropyrimidine inhibitors of Rho kinase (ROCK1) as potential therapeutic agents for cardiovascular diseases [J]. J Med Chem. 2008 Nov 13;51(21):6631-4.

Yuan Y, Xu J, Jiang L, et al. Discovery, Optimization, and Structure-Activity Relationship Study of Novel and Potent RSK4 Inhibitors as Promising Agents for the Treatment of Esophageal Squamous Cell Carcinoma. Journal of Medicinal Chemistry. 2021

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