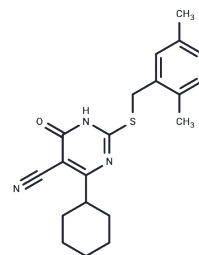


ESI-08

Chemical Properties

CAS No. : 301177-43-5
 Formula: C₂₀H₂₃N₃O₅
 Molecular Weight: 353.48
 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	ESI-08 is an effective antagonist of EPAC2 with an IC ₅₀ of 8.4 μM. ESI-08 selectively blocks cAMP-induced EPAC activation but not cAMP-mediated PKA activation.
Targets(IC ₅₀)	cAMP,Ras
In vitro	ESI-08 (25 μM) does not alter cAMP-induced type I and II PKA holoenzymes activation while H89, a selective PKA inhibitor, blocked the type I or II PKA activities completely[1].

Solubility Information

Solubility	DMSO: 50 mg/mL (141.45 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.829 mL	14.1451 mL	28.2901 mL
5 mM	0.5658 mL	2.829 mL	5.658 mL
10 mM	0.2829 mL	1.4145 mL	2.829 mL
50 mM	0.0566 mL	0.2829 mL	0.5658 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

Chen H, et al. 5-Cyano-6-oxo-1,6-dihydro-pyrimidines as potent antagonists targeting exchange proteins directly activated by cAMP. *Bioorg Med Chem Lett.* 2012 Jun 15;22(12):4038-43.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

This product is for Research Use Only · Not for Human or Veterinary or Therapeutic Use

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