Data Sheet (Cat.No.T2415)



PP121

Chemical Properties

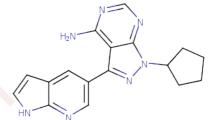
CAS No.: 1092788-83-4

Formula: C17H17N7

Molecular Weight: 319.36

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year



Biological Description

| Description | PP-121 is a multi-targeted inhibitor of PDGFR (IC50: 2 nM), Hck (IC50: 8 nM), mTOR (IC50: 10 nM), VEGFR2(IC50: 12 nM), Src (IC50: 14 nM) and Abl (IC50: 18 nM), also inhibits DNA-PK (IC50: 60 nM). |
|---------------|---|
| Targets(IC50) | Apoptosis,VEGFR,Bcr-Abl,PDGFR,Src,mTOR,Hck |
| Kinase Assay | Kinase assays: Purified kinase domains are incubated with PP-121 at 2- or 4-fold dilutions over a concentration range of 1 nM-50 μ M or with vehicle (0.1% DMSO) in the presence of 10 μ M ATP, 2.5 μ Ci of γ -32P-ATP and substrate. Reactions are terminated by spotting onto nitrocellulose or phosphocellulose membranes, depending on the substrate; this membrane is then washed 5-6 times to remove unbound radioactivity and dried. Transferred radioactivity is quantitated by phosphorimaging and IC50 values are calculated by fitting the data to a sigmoidal doseresponse using Prism software. |
| Cell Research | For western blot analysis, cells are grown in 12-well plates and treated with PP-121 at the indicated concentrations or vehicle (0.1% DMSO). Treated cells are lysed, lysates are resolved by SDS-PAGE, transferred to nitrocellulose and blotted. For cell proliferation assays, cells are grown in 96-well plates are treated with PP-121 at 4-fold dilutions (10 µM - 0.040 µM) or vehicle (0.1% DMSO). After 72 hours cells are exposed to Resazurin sodium salt (22 µM) and fluorescence is quantified. IC50 values are calculated using Prism software. For proliferation assays involving single cell counting, non-adherent cells are plated at low density (3-5% confluence) and treated with PP-121 (2.5 µM) or vehicle (0.1% DMSO). Cells are diluted into trypan blue daily and viable cells counted using a hemocytometer. For apoptosis and cell cycle analysis, cells are treated with the indicated concentration of PP-121 or vehicle (0.1% DMSO) for 24-72 hours. Cells are either stained live with AnnexinV-FITC or fixed with ethanol and stained with propidium iodide. Cell populations are separated using a FacsCalibur flow cytometer; data is collected using CellQuest Pro software and analyzed with either ModFit or FlowJo Software.(Only for Reference) |

Solubility Information

| Solubility | H2O: < 1 mg/mL (insoluble or slightly soluble), Ethanol: 2 mg/mL (6.26 |
|------------|---|
| | mM), DMSO: 60 mg/mL (187.9 mM), (< 1 mg/ml refers to the product |
| .0, | slightly soluble or insoluble) |

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Preparing Stock Solutions

| | 1mg | 5mg | 10mg |
|-------|-----------|------------|------------|
| 1 mM | 3.1313 mL | 15.6563 mL | 31.3126 mL |
| 5 mM | 0.6263 mL | 3.1313 mL | 6.2625 mL |
| 10 mM | 0.3131 mL | 1.5656 mL | 3.1313 mL |
| 50 mM | 0.0626 mL | 0.3131 mL | 0.6263 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

Zhou D, Yang S, Yan H, et al. SC75741, a novel c-Abl inhibitor, promotes the clearance of TDP25 aggregates via ATG5-dependent autophagy pathway. Frontiers in Pharmacology. 2021: 2891.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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