

PKI-402

Chemical Properties

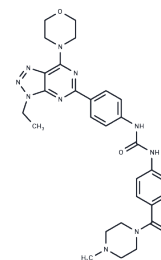
CAS No. : 1173204-81-3

Formula: C₂₉H₃₄N₁₀O₃

Molecular Weight: 570.65

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 | PKI-402 is a potent PI3K and mTOR inhibitor. PKI-402 inhibits PI3K α , mTOR, PI3K β , PI3K δ and PI3K γ with IC ₅₀ s of 2, 3, 7, 14 and 16 nM, respectively. |
| Targets(IC ₅₀) | mTOR,PI3K |
| In vitro | PKI-402 causes in vitro growth inhibition of human tumor cell lines derived from a diverse set of human tumor tissues, including breast, brain (glioma), pancreas, and non-small cell lung cancer (NSCLC) tissues[1]. |
| In vivo | PKI-402 exhibits antitumor efficacy via intravenous administration in breast [MDA-MB-361: Her2+ and PIK3CA (E545K)], glioma (U87MG and PTEN), and NSCLC (A549; K-Ras and STK11) xenograft models, with induced regression observed specifically in the MDA-MB-361 model. |

Solubility Information

| | |
|---------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Solubility | DMSO: 4.68 mg/mL (8.2 mM),Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble) |
| In vivo Formulation | 10% DMSO+90% Saline: 0.47 mg/mL (0.82 mM),Suspension. <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 | 1.7524 mL | 8.7619 mL | 17.5239 mL |
| 5 mM | 0.3505 mL | 1.7524 mL | 3.5048 mL |
| 10 mM | 0.1752 mL | 0.8762 mL | 1.7524 mL |
| 50 mM | 0.035 mL | 0.1752 mL | 0.3505 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

Mallon R et al. Antitumor efficacy profile of PKI-402, a dual phosphatidylinositol 3-kinase/mammalian target of rapamycin inhibitor. Mol Cancer Ther. 2010 Apr;9(4):976-84.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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