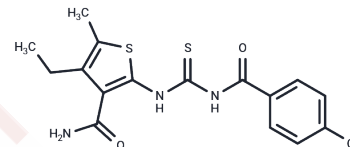


PI-273

## Chemical Properties

CAS No. : 925069-34-7  
 Formula: C<sub>16</sub>H<sub>16</sub>ClN<sub>3</sub>O<sub>2</sub>S<sub>2</sub>  
 Molecular Weight: 381.9  
 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	PI-273, a Substrate-Competitive, Specific Small-Molecule Inhibitor of PI4KII $\alpha$ , Inhibits the Growth of Breast Cancer Cells
Targets(IC50)	Apoptosis,PI4K
In vitro	PI-273 exhibited the greatest inhibitory effect on PI4KII $\alpha$ kinase activity (IC <sub>50</sub> = 0.47 $\mu$ mol/L) and suppressed cell proliferation. Surface plasmon resonance and thermal shift assays indicated that PI-273 interacted directly with PI4KII $\alpha$ . The kinetic analysis identified PI-273 as a reversible competitive inhibitor with respect to the substrate phosphatidylinositol (PI), which contrasted with most other PI kinase inhibitors that bind the ATP binding site. PI-273 reduced PI4P content, cell viability, and AKT signaling in wild-type MCF-7 cells, but not in PI4KII $\alpha$ knockout MCF-7 cells, indicating that PI-273 is highly selective for PI4KII $\alpha$ . Mutant analysis revealed the role of palmitoylation insertion in the selectivity of PI-273 for PI4KII $\alpha$ . In addition, PI-273 treatment retarded cell proliferation by blocking cells in G <sub>2</sub> -M, inducing cell apoptosis and suppressing colony-forming ability. Importantly, PI-273 significantly inhibited MCF-7 cell-induced breast tumor growth without toxicity. PI-273 is the first substrate-competitive, a subtype-specific inhibitor of PI4KII $\alpha$ , the use of which will facilitate evaluations of PI4KII $\alpha$ as a cancer therapeutic target.
In vivo	PI-273 (intraperitoneal injection; 25 mg/kg/day; 15 days) significantly reduces tumor volume and weight in MCF-7 xenografts [1]. PI-273 (0.5 mg/kg intravenously or 1.5 mg/kg intragastrically; 0.08-5 hours) has a half-life of 0.411 hours for intravenous administration and 1.321 hours for intragastric administration, with an absolute bioavailability of 5.1% [1].

## Solubility Information

Solubility DMSO: 0.95 mg/mL (2.49 mM),Sonication is recommended.  
 (< 1 mg/ml refers to the product slightly soluble or insoluble)

### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.6185 mL	13.0924 mL	26.1849 mL
5 mM	0.5237 mL	2.6185 mL	5.237 mL
10 mM	0.2618 mL	1.3092 mL	2.6185 mL
50 mM	0.0524 mL	0.2618 mL	0.5237 mL

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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

Li J, et al. PI-273, a Substrate-Competitive, Specific Small-Molecule Inhibitor of PI4KII $\alpha$ , Inhibits the Growth of Breast Cancer Cells. *Cancer Res.* 2017 Nov 15;77(22):6253-6266.

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