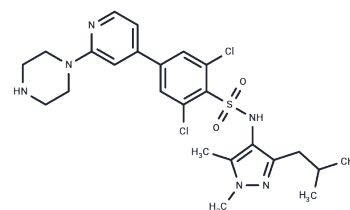


PCLX-001

## Chemical Properties

CAS No. : 1215011-08-7  
Formula: C<sub>24</sub>H<sub>30</sub>Cl<sub>2</sub>N<sub>6</sub>O<sub>2</sub>S  
Molecular Weight: 537.51  
Appearance: no data available  
Storage: store at low temperature, keep away from direct sunlight  
Powder: -20°C for 3 years | In solvent: -80°C for 1 year



## Biological Description

|                            |                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              |
|----------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Description                | PCLX-001 is a small-molecule compound that acts as an orally active inhibitor of N-myristoyltransferase (NMT), specifically targeting NMT1 and NMT2 with IC <sub>50</sub> values of 5 nM and 8 nM, respectively. This compound demonstrates anti-tumor effects and effectively inhibits the early signaling of B-cell receptor (BCR). Consequently, PCLX-001 is a valuable tool for researching B-cell malignancies [1] [2].                                                                                                                                                                 |
| Targets(IC <sub>50</sub> ) | Others                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |
| In vitro                   | We discovered a marked sensitivity of hematological cancer cell lines, including B-cell lymphomas, to the potent pan-NMT inhibitor PCLX-001. PCLX-001 treatment impacts the global myristoylation of lymphoma cell proteins and inhibits early B-cell receptor (BCR) signaling events critical for survival. In addition to abrogating myristoylation of Src family kinases, PCLX-001 also promotes their degradation and, unexpectedly, that of numerous non-myristoylated BCR effectors including c-Myc, NFκB and P-ERK, leading to cancer cell death in vitro and in xenograft models.[2] |
| In vivo                    | Treatment of cultured breast cancer cells with PCLX-001 reduced cell viability in vitro. Daily oral administration of PCLX-001 to immunodeficient mice bearing human MDA-MB-231 breast cancer xenografts produced significant dose-dependent tumor growth inhibition in vivo.[4]                                                                                                                                                                                                                                                                                                             |

## Solubility Information

|            |                                                                                             |
|------------|---------------------------------------------------------------------------------------------|
| Solubility | DMSO: 5.38 mg/mL (10 mM)<br>(< 1 mg/ml refers to the product slightly soluble or insoluble) |
|------------|---------------------------------------------------------------------------------------------|

## A DRUG SCREENING EXPERT

### Preparing Stock Solutions

|       | 1mg       | 5mg       | 10mg       |
|-------|-----------|-----------|------------|
| 1 mM  | 1.8604 mL | 9.3022 mL | 18.6043 mL |
| 5 mM  | 0.3721 mL | 1.8604 mL | 3.7209 mL  |
| 10 mM | 0.186 mL  | 0.9302 mL | 1.8604 mL  |
| 50 mM | 0.0372 mL | 0.186 mL  | 0.3721 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

Weickert M, et al. Initial characterization and toxicology of an nmt inhibitor in development for hematologic malignancies. Blood. 2019;134: 3362.

**Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins**

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