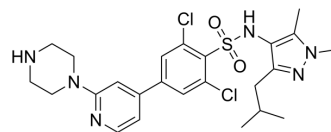


PCLX-001

Cat. No.:	HY-147308		
CAS No.:	1215011-08-7		
Molecular Formula:	C ₂₄ H ₃₀ Cl ₂ N ₆ O ₂ S		
Molecular Weight:	537.51		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : 10 mg/mL (18.60 mM; ultrasonic and warming and heat to 60°C)

Concentration	Solvent	Mass	1 mg			5 mg			10 mg		
			Concentration			Concentration			Concentration		
1 mM			1.8604 mL			9.3022 mL			18.6043 mL		
5 mM			0.3721 mL			1.8604 mL			3.7209 mL		
10 mM			0.1860 mL			0.9302 mL			1.8604 mL		

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 1 mg/mL (1.86 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 1 mg/mL (1.86 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 1 mg/mL (1.86 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

PCLX-001 is an orally active, small-molecule, dual N-myristoyltransferase (NMT) inhibitor, with IC₅₀s of 5 nM (NMT1) and 8 nM (NMT2), respectively. PCLX-001 exhibits anti-tumor activity and inhibits early B-cell receptor (BCR) signaling, can be used to B-cell malignancies research^{[1][2]}.

REFERENCES

[1]. Michael Weickert, et al. Initial Characterization and Toxicology of an Nmt Inhibitor in Development for Hematologic Malignancies. Blood. 2019. 134(s1):3362.

[2]. Beauchamp E, et al. Targeting N-myristoylation for therapy of B-cell lymphomas. Nat Commun. 2020 Oct 22;11(1):5348.

Caution: Product has not been fully validated for medical applications. For research use only.

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