Product Data Sheet

LY-3381916

Cat. No.: HY-111540 CAS No.: 2166616-75-5 Molecular Formula: $C_{23}H_{25}FN_{2}O_{3}$ Molecular Weight: 396.45

Target: Indoleamine 2,3-Dioxygenase (IDO)

Pathway: Metabolic Enzyme/Protease Powder

3 years 4°C 2 years

In solvent -80°C 2 years

-20°C

-20°C 1 year

SOLVENT & SOLUBILITY

In Vitro

Storage:

DMSO: 300 mg/mL (756.72 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.5224 mL	12.6119 mL	25.2239 mL
	5 mM	0.5045 mL	2.5224 mL	5.0448 mL
	10 mM	0.2522 mL	1.2612 mL	2.5224 mL

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

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (6.31 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	LY-3381916 (IDO1-IN-5) is a potent, selective and brain penetrated inhibitor of IDO1 activity, binds to apo-IDO1 lacking heme rather than mature heme-bound IDO1 $^{[1]}$.	
IC ₅₀ & Target	IDO1	
In Vitro	LY-3381916 (IDO1-IN-5) (up to 100 μ M) shows no obvious agonism of aryl hydrocarbon receptor (AHR) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

CUSTOMER VALIDATION

- Immune Netw. 2020 Oct 19;20(5):e36.
- The University of Auckland. 2023 Mar.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Frank C. Dorsey, et al. Abstract 5245: Identification and characterization of the IDO1 inhibitor LY3381916. Cancer Research. 2018, 78(13).

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

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