Inhibitors

IDO-IN-8

Cat. No.: HY-18770C CAS No.: 1402837-77-7 Molecular Formula: $\mathsf{C}_{18}\mathsf{H}_{21}\mathsf{FN}_2\mathsf{O}_2$ Molecular Weight: 316.37

Target: Indoleamine 2,3-Dioxygenase (IDO)

Pathway: Metabolic Enzyme/Protease

> Powder -20°C 3 years $4^{\circ}C$ 2 years

> -80°C In solvent 2 years

> > -20°C 1 year

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

Storage:

Ethanol: 100 mg/mL (316.09 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.1609 mL	15.8043 mL	31.6086 mL
	5 mM	0.6322 mL	3.1609 mL	6.3217 mL
	10 mM	0.3161 mL	1.5804 mL	3.1609 mL

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

In Vivo

- 1. Add each solvent one by one: 10% EtOH >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (7.90 mM); Clear solution
- 2. Add each solvent one by one: 10% EtOH >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.90 mM); Clear solution
- 3. Add each solvent one by one: 10% EtOH >> 90% corn oil Solubility: ≥ 2.5 mg/mL (7.90 mM); Clear solution

BIOLOGICAL ACTIVITY

Description IDO-IN-8 (NLG-1487) is an indoleamine 2,3-dioxygenase (IDO) inhibitor extracted from patent WO WO2012142237A1, compound 1487, has an IC $_{50}$ of 1-10 μ M.

IC₅₀ & Target IDO

 $1-10~\mu M~(IC_{50})$

In Vitro IDO-IN-8 (Compound 1487) is an indoleamine 2,3-dioxygenase (IDO) inhibitor with an IC $_{50}$ of 1-10 μ M (this is the

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	concentration of IDO-IN-8 at which inhibits 50% of enzymatic activity using recombinant human IDO) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
REFERENCES		
1]. Mautino, et al. Preparati	on of fused imidazole derivatives as IDO inhibitors. From PCT Int. Appl. (2012), 20121018.	

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

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