## **Product** Data Sheet

## **Povorcitinib**

 Cat. No.:
 HY-145588 

 CAS No.:
 1637677-22-5 

 Molecular Formula:
  $C_{23}H_{22}F_5N_7O$  

 Molecular Weight:
 507.46 

Pathway: Epigenetics; JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Stem Cell/Wnt

Storage: Powder -20°C 3 years 4°C 2 years

JAK

In solvent  $-80^{\circ}$ C 6 months  $-20^{\circ}$ C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

Target:

DMSO: 250 mg/mL (492.65 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9706 mL	9.8530 mL	19.7060 mL
	5 mM	0.3941 mL	1.9706 mL	3.9412 mL
	10 mM	0.1971 mL	0.9853 mL	1.9706 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: ≥ 2.08 mg/mL (4.10 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.08 mg/mL (4.10 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.10 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Povorcitinib (INCB54707) is a potent and selective JAK1 inhibitor that effectively reduces abscesses and inflammatory nodules. Povorcitinib is used to study cutaneous lupus erythematosus (CLE) and lichen planus (LP) (information extracted from patent WO2021076124A1).
IC <sub>50</sub> & Target	$JAK1^{[1]}$

REFERENCES				
[1]. Paul Smith, et al. Use of jak1 inhibitors for the treatment of cutaneous lupus erythematosus and lichen planus (lp). Patent WO2021076124A1.				
Caution: Product has not been fully validated for mo				
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