Product Data Sheet

FLT3/CHK1-IN-1

Cat. No.: HY-155195 CAS No.: 2991054-23-8 Molecular Formula: $C_{25}H_{33}F_3N_6O_2$

Molecular Weight: 506.56 Others Target: Pathway: Others

4°C, protect from light Storage:

* In solvent: -80°C, 6 months; -20°C, 1 month (protect from light)

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (197.41 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9741 mL	9.8705 mL	19.7410 mL
	5 mM	0.3948 mL	1.9741 mL	3.9482 mL
	10 mM	0.1974 mL	0.9870 mL	1.9741 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 (4.94 mM); Clear solution; Need ultrasonic
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (4.94 mM); Clear solution; Need ultrasonic
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (4.94 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description

FLT3/CHK-IN-1 (Compound 18) is a dual inhibitor of FLT3/CHK1. FLT3/CHK-IN-1 is more than 1700 times more selective to c-KIT and greatly reduces hERG affinity with an IC $_{50}$ value of 58.4 μ M. FLT3/CHK-IN-1 inhibits tumor growth in mouse xenotransplantation models inoculated with MV-4-11 cells $^{[1]}$.

REFERENCES

[1]. Li X, et al. Discovery of 2-Aminopyrimidine Derivatives as Potent Dual FLT3/CHK1 Inhibitors with Significantly Reduced hERG Inhibitory Activities. J Med Chem. 2023 Sep.

14;66(17):11792-11814.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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