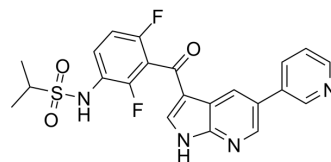


BRAF inhibitor

Cat. No.:	HY-10247		
CAS No.:	918505-61-0		
Molecular Formula:	C ₂₂ H ₁₈ F ₂ N ₄ O ₃ S		
Molecular Weight:	456.47		
Target:	Raf		
Pathway:	MAPK/ERK Pathway		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro

DMSO : 50 mg/mL (109.54 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.1907 mL	10.9536 mL	21.9072 mL
	5 mM	0.4381 mL	2.1907 mL	4.3814 mL
	10 mM	0.2191 mL	1.0954 mL	2.1907 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: ≥ 2.5 mg/mL (5.48 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 2.5 mg/mL (5.48 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 2.5 mg/mL (5.48 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

BRAF inhibitor is a B-Raf inhibitor extracted from patent WO/2011103196 A1, Compound P-0850.

IC₅₀ & Target

B-Raf

CUSTOMER VALIDATION

-
- Cell Death Dis. 2022 Jul 15;13(7):615.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Prahbha N. Ibrahim, et al. Pyrrolo[2,3-b] pyridine derivatives as protein kinase inhibitors. WO2007002325A1.

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

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