Exarafenib

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway:	HY-147268 2639957-39-2 C ₂₆ H ₃₄ F ₃ N ₅ O ₃ 521.58 Raf; p38 MAPK MAPK/ERK Pathway	
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)	

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (191.73 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	1.9173 mL	9.5863 mL	19.1725 mL	
		5 mM	0.3835 mL	1.9173 mL	3.8345 mL	
		10 mM	0.1917 mL	0.9586 mL	1.9173 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 (4.79 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (4.79 mM); Suspended solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description Exarafenib (RAF/KIN_2787) is a potent, orally active pan-RAF inhibitor. Exarafenib has anticancer activity by suppression of downstream MAPK pathway signaling. Exarafenib can be used for cancer research^{[1][2]}.

REFERENCES

[1]. Miller N, et, al. Antitumor activity of KIN-2787, a next-generation pan-RAF inhibitor, in combination with MEK inhibition in preclinical models of human NRAS mutant melanoma.2022 Jun 2;40(16): e15099.

[2]. WHO Drug Information. International Nonproprietary Names for Pharmaceutical

Product Data Sheet



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

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