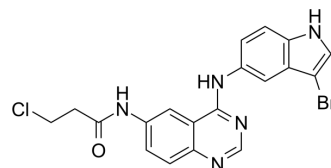


pan-HER-IN-2

Cat. No.:	HY-144677		
CAS No.:	1639040-95-1		
Molecular Formula:	C ₁₉ H ₁₅ BrClN ₅ O		
Molecular Weight:	444.71		
Target:	EGFR; Apoptosis		
Pathway:	JAK/STAT Signaling; Protein Tyrosine Kinase/RTK; Apoptosis		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (224.87 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.2487 mL	11.2433 mL	22.4866 mL
		5 mM	0.4497 mL	2.2487 mL	4.4973 mL
10 mM		0.2249 mL	1.1243 mL	2.2487 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 (5.62 mM); Clear solution; Need ultrasonic				

BIOLOGICAL ACTIVITY

Description	pan-HER-IN-2 (Compound C6) is a reversible, orally active pan-HER inhibitor with IC ₅₀ values of 0.72, 2.0, 8.2 and 75.1 nM against EGFR, HER4, EGFR ^{T790M/L858R} and HER2, respectively. pan-HER-IN-2 induces apoptosis and shows antitumor activities ^[1] .			
IC₅₀ & Target	EGFR 0.72 nM (IC ₅₀)	HER4 2.0 nM (IC ₅₀)	EGFR (L858R/T790M) 8.2 nM (IC ₅₀)	HER2 75.1 nM (IC ₅₀)

REFERENCES

[1]. Tang Q, et al. Discovery of N-(3-bromo-1H-indol-5-yl)-quinazolin-4-amine as an effective molecular skeleton to develop reversible/irreversible pan-HER inhibitors. Eur J

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

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