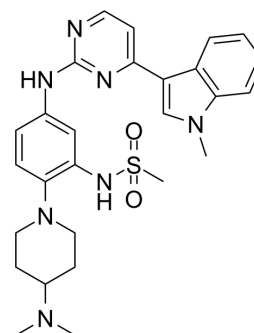


Os30

Cat. No.:	HY-155358		
CAS No.:	2998928-68-8		
Molecular Formula:	C ₂₇ H ₃₃ N ₇ O ₂ S		
Molecular Weight:	519.66		
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 : 50 mg/mL (96.22 mM); ultrasonic and warming and heat to 80°C)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.9243 mL	9.6217 mL	19.2434 mL
5 mM	0.3849 mL	1.9243 mL	3.8487 mL
10 mM	0.1924 mL	0.9622 mL	1.9243 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 (4.81 mM); Clear solution; Need ultrasonic

BIOLOGICAL ACTIVITY

Description

Os30, a potent fourth-generation EGFR inhibitor, is a potent EGFR C797S-TK inhibitor with IC₅₀ values of 18 nM and 113 nM for EGFR Del19/T790M/C797S TK and EGFR L858R/T790M/C797S TK, respectively. Os30 can suppress EGFR phosphorylation, arrest at G1 phase and induce the apoptosis of KC-0116 (BaF3-EGFR Del19/T790M/C797S) cells. Os30 shows potent antitumor efficacy on non-small cell lung cancer (NSCLC) with EGFR C797S mutation^[1].

REFERENCES

[1]. Xiao-Xiao Xi, et al. Modification of osimertinib to discover new potent EGFR C797S-TK inhibitors. Eur J Med Chem. 2023 Dec 5;261:115865.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA