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Zuschläge

- Mindermengenzuschlag
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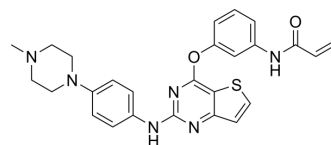
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Olmudinib

Cat. No.:	HY-19730		
CAS No.:	1353550-13-6		
Molecular Formula:	C ₂₆ H ₂₆ N ₆ O ₂ S		
Molecular Weight:	486.59		
Target:	EGFR		
Pathway:	JAK/STAT Signaling; Protein Tyrosine Kinase/RTK		
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 : 125 mg/mL (256.89 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		2.0551 mL	10.2756 mL	20.5512 mL
		5 mM		0.4110 mL	2.0551 mL	4.1102 mL
		10 mM		0.2055 mL	1.0276 mL	2.0551 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: ≥ 6.25 mg/mL (12.84 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 6.25 mg/mL (12.84 mM); Clear solution; Need ultrasonic					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 6.25 mg/mL (12.84 mM); Suspended solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description	Olmudinib (HM61713; BI-1482694) is an orally active and irreversible third EGFR tyrosine kinase inhibitor that binds to a cysteine residue near the kinase domain. Olmutinib is used for NSCLC ^{[1][2]} .	
IC ₅₀ & Target	EGFR ^{Exon 19 deletion} 9.2 nM (IC ₅₀ , Cell Assay)	EGFR ^{L858R/T790M} 10 nM (IC ₅₀ , Cell Assay)
In Vitro	Olmudinib potently inhibits EGFR in HCC827 cells expressing EGFR ^{DEL19} (IC ₅₀ =9.2 nM) and H1975 cells expressing EGFR	

L858R/T790M ($IC_{50}=10$ nM). In contrast, the IC_{50} of olmutinib against cells expressing EGFR^{WT} is 2225 nM^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Acta Pharm Sin B. 2018 Jul;8(4):563-574.
- Mol Syst Biol. 2023 Dec 18.
- J Pharm Anal. 2021 Jun 19.
- Biomedicines. 2024 Jan 7, 12(1), 123.
- J Biomol Struct Dyn. 2021 Nov 12;1-11.

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REFERENCES

[1]. Kim ES, et al. Olmutinib: First Global Approval. Drugs. 2016 Jul;76(11):1153-7.

[2]. Mohamed W. Attwa, et al. Detection and characterization of olmutinib reactive metabolites by LC-MS/MS: Elucidation of bioactivation pathways. Journal of Separation science. 18 November 2019.

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

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