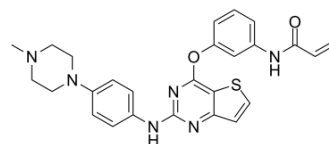


## Olmutinib

<b>Cat. No.:</b>	HY-19730		
<b>CAS No.:</b>	1353550-13-6		
<b>Molecular Formula:</b>	C <sub>26</sub> H <sub>26</sub> N <sub>6</sub> O <sub>2</sub> S		
<b>Molecular Weight:</b>	486.59		
<b>Target:</b>	EGFR		
<b>Pathway:</b>	JAK/STAT Signaling; Protein Tyrosine Kinase/RTK		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : ≥ 44 mg/mL (90.43 mM)  
 \* "≥" means soluble, but saturation unknown.

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

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
 Solubility: ≥ 2.08 mg/mL (4.27 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
 Solubility: ≥ 2.08 mg/mL (4.27 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
 Solubility: ≥ 2.08 mg/mL (4.27 mM); Clear solution

### BIOLOGICAL ACTIVITY

#### Description

Olmutinib (HM61713; BI-1482694) is an orally bioavailable and irreversible third EGFR tyrosine kinase inhibitor that binds to a cysteine residue near the kinase domain. Olmutinib is used for NSCLC<sup>[1][2]</sup>.

#### IC<sub>50</sub> & Target

EGFR <sup>Exon 19 deletion</sup>	EGFR <sup>L858R/T790M</sup>
9.2 nM (IC <sub>50</sub> , Cell Assay)	10 nM (IC <sub>50</sub> , Cell Assay)

## In Vitro

Olmudinib potently inhibits EGFR in HCC827 cells expressing EGFR<sup>DEL19</sup> (IC<sub>50</sub>=9.2 nM) and H1975 cells expressing EGFR L858R/T790M (IC<sub>50</sub>=10 nM). In contrast, the IC<sub>50</sub> of olmutinib against cells expressing EGFR<sup>WT</sup> is 2225 nM<sup>[1]</sup>. 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.
- RSC Adv. 2018, 8, 40387-40394.
- J Sep Sci. 2020 Feb;43(4):708-718.
- Med Sci Monit. 2020 Aug 24;26:e924922.

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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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