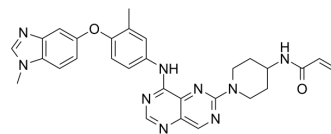


## Zongertinib

Cat. No.:	HY-148810		
CAS No.:	2728667-27-2		
Molecular Formula:	C <sub>29</sub> H <sub>29</sub> N <sub>9</sub> O <sub>2</sub>		
Molecular Weight:	535.6		
Target:	c-Met/HGFR		
Pathway:	Protein Tyrosine Kinase/RTK		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

In Vitro	DMSO : 31.25 mg/mL (58.35 mM; ultrasonic and warming and heat to 60°C)					
	Preparing Stock Solutions	Solvent Concentration	Mass			
			1 mg	5 mg	10 mg	
			1 mM	1.8671 mL	9.3353 mL	18.6707 mL
			5 mM	0.3734 mL	1.8671 mL	3.7341 mL
10 mM	0.1867 mL	0.9335 mL	1.8671 mL			
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.67 mM); Clear solution					

### BIOLOGICAL ACTIVITY

Description	Zongertinib (BI 1810631) is a potent and selective HER2 and EGFR tyrosine kinase inhibitor with IC <sub>50</sub> values of 13 nM and 579 nM, respectively. Zongertinib has antitumor activity and can be used in the study of multiple solid tumors <sup>[1][2][3]</sup> .
In Vitro	Zongertinib in NCI-H-2170 HER2 wt amp, NCI-H2170 HER2 YVMA, A431 EGFR wt amp, BAF3 HER2 WT, BAF3 HER2 YVMA and BAF3 EGF dep. The IC <sub>50</sub> values that inhibits cell proliferation are 6 nM, 33 nM, >5000 nM, 1 nM, 16 nM, and 1540 nM, respectively <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. WHO Drug Information - World Health Organization (WHO).

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[2]. Wilding Birgit, et al. Synthesis of diazino-pyrimidines as anticancer agents: World Intellectual Property Organization, WO2021213800. 2021-10-28.

[3]. Li S, et al. Emerging Targeted Therapies in Advanced Non-Small-Cell Lung Cancer. *Cancers (Basel)*. 2023 May 24;15(11):2899.

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

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