**Proteins** 

# **Almonertinib**

Cat. No.: HY-112823 CAS No.: 1899921-05-1 Molecular Formula:  $C_{30}H_{35}N_{7}O_{2}$ Molecular Weight: 525.64 **EGFR** Target:

Pathway: JAK/STAT Signaling; Protein Tyrosine Kinase/RTK

-20°C Storage: Powder 3 years

 $4^{\circ}C$ 2 years

-80°C In solvent 2 years

> -20°C 1 year

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**Product** Data Sheet

### **SOLVENT & SOLUBILITY**

In Vitro DMSO:  $\geq$  83.33 mg/mL (158.53 mM)

\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9024 mL	9.5122 mL	19.0244 mL
	5 mM	0.3805 mL	1.9024 mL	3.8049 mL
	10 mM	0.1902 mL	0.9512 mL	1.9024 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 (11.89 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 6.25 mg/mL (11.89 mM); Clear solution

### **BIOLOGICAL ACTIVITY**

Almonertinib (HS-10296) is an orally available, irreversible, third-generation EGFR tyrosine kinase inhibitor with high Description

> selectivity for EGFR-sensitizing and T790M resistance mutations. Almonertinib shows great inhibitory activity against T790M, T790M/L858R and T790M/Del19 (IC<sub>50</sub>: 0.37, 0.29 and 0.21 nM, respectively), and is less effective against wild type (3.39 nM).

Almonertinib is used for the research of the non-small cell lung cancer [1][2].

EGFR<sup>L858R</sup>/T790M EGFR<sup>T790M</sup> EGFR<sup>del19</sup> T790M IC<sub>50</sub> & Target

0.37 nM (IC<sub>50</sub>) 0.29 nM (IC<sub>50</sub>) 0.21 nM (IC<sub>50</sub>)

In Vitro HS-10296 is an orally available inhibitor of the epidermal growth factor receptor (EGFR) mutant form T790M, with potential antineoplastic activity, which canbe used to treat  $NSCLC^{[2]}$ . Additionaly, HS-10296 could also inhibit other EGFR sensitive mutations, including G719X, del19, L858R and L861Q<sup>[3]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **CUSTOMER VALIDATION**

- Cell Rep Med. 2023 Jan 10;100911.
- Front Pharmacol. 2021 May 14;12:671328.
- Patent. US20220177473A1.

See more customer validations on www.MedChemExpress.com

#### **REFERENCES**

- [1]. Sullivan I, et al. Next-Generation EGFR Tyrosine Kinase Inhibitors for Treating EGFR-Mutant Lung Cancer beyond First Line. Front Med (Lausanne). 2017 Jan 18;3:76.
- [2]. Wu SG, et al. Management of acquired resistance to EGFR TKI-targeted therapy in advanced non-small cell lung cancer. Mol Cancer. 2018 Feb 19;17(1):38.
- [3]. Yang JC, et al. Safety, Efficacy, and Pharmacokinetics of Almonertinib (HS-10296) in Pretreated Patients With EGFR-Mutated Advanced NSCLC: A Multicenter, Openlabel, Phase 1 Trial [published online ahead of print, 2020 Sep 9]. J Thorac Oncol. 2020;S1556-0

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

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