

## **Product** Data Sheet

## **Enbezotinib**

Cat. No.:HY-145565CAS No.:2359649-81-1Molecular Formula: $C_{21}H_{21}FN_6O_3$ Molecular Weight:424.43Target:RET

Pathway: Protein Tyrosine Kinase/RTK

Storage: Powder -20°C 3 years

 $\begin{tabular}{ll} $4^{\circ}C$ & 2 years \\ $\text{In solvent}$ & $-80^{\circ}C$ & 6 months \\ \end{tabular}$ 

-20°C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 100 mg/mL (235.61 mM; ultrasonic and warming and heat to 80°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3561 mL	11.7805 mL	23.5610 mL
	5 mM	0.4712 mL	2.3561 mL	4.7122 mL
	10 mM	0.2356 mL	1.1781 mL	2.3561 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: ≥ 2.08 mg/mL (4.90 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.90 mM); Clear solution

## **BIOLOGICAL ACTIVITY**

Description	Enbezotinib, an inhibitor of RET, can inhibit the RET autophosphorylation. Enbezotinib can be used for the research of cancer $^{[1]}$ .
IC <sub>50</sub> & Target	$RET^{[1]}$
In Vitro	Enbezotinib (compound 5) (0.3-300 nM) inhibits p-RET (y905) in RET-driven cells <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	Enbezotinib (compound 5) (2-5 mg/kg; twice daily for 27 days) decreases tumor size in $mice^{[1]}$ . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES							
[1]. ROGERS EW, et, al. Macrocyclic compounds for treating disease. WO2019126121A1.							
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