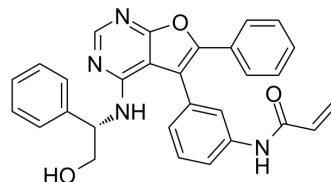


## EGFR-IN-9

<b>Cat. No.:</b>	HY-18213		
<b>CAS No.:</b>	1226549-39-8		
<b>Molecular Formula:</b>	C <sub>29</sub> H <sub>24</sub> N <sub>4</sub> O <sub>3</sub>		
<b>Molecular Weight:</b>	476.53		
<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 : 100 mg/mL (209.85 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	2.0985 mL	10.4925 mL	20.9850 mL
5 mM	0.4197 mL	2.0985 mL	4.1970 mL
10 mM	0.2099 mL	1.0493 mL	2.0985 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

<b>Description</b>	EGFR-IN-9 (Compound 8) is a potent EGFR kinase inhibitor with IC <sub>50</sub> s of 7 nM, 28 nM for the wild type EGFR kinase and double mutant EGFR kinase (L858R/T790M). EGFR-IN-9 has antitumor activity <sup>[1]</sup> .	
<b>IC<sub>50</sub> &amp; Target</b>	EGFR (WT) 7 nM (IC <sub>50</sub> )	EGFR <sup>L858R/T790M</sup> 28 nM (IC <sub>50</sub> )
<b>In Vitro</b>	EGFR-IN-9 (Compound 8) shows antiproliferative activity in HCC827 lung cancer cell line with an IC <sub>50</sub> of 8 nM. MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

### REFERENCES

[1]. Coumar MS, et al. Fast-forwarding hit to lead: aurora and epidermal growth factor receptor kinase inhibitor leadidentification. J Med Chem. 2010 Jul 8;53(13):4980-8.

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**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](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA