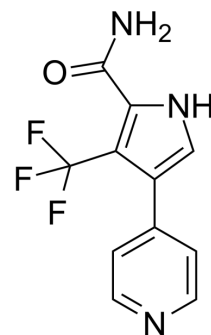


## CDK8-IN-1

<b>Cat. No.:</b>	HY-103492		
<b>CAS No.:</b>	1629633-48-2		
<b>Molecular Formula:</b>	C <sub>11</sub> H <sub>8</sub> F <sub>3</sub> N <sub>3</sub> O		
<b>Molecular Weight:</b>	255.2		
<b>Target:</b>	CDK		
<b>Pathway:</b>	Cell Cycle/DNA Damage		
<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 (391.85 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	3.9185 mL	19.5925 mL	39.1850 mL
5 mM	0.7837 mL	3.9185 mL	7.8370 mL
10 mM	0.3918 mL	1.9592 mL	3.9185 mL

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

### BIOLOGICAL ACTIVITY

#### Description

CDK8-IN-1 is a potent and selective CDK8 inhibitor with an IC<sub>50</sub> of 3 nM.

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

CDK8  
3 nM (IC<sub>50</sub>)

#### In Vivo

CDK8-IN-1 displays low systemic clearance, very good exposure and oral bioavailability. The t<sub>max</sub> is 0.25 h by PO. The mean values of C<sub>max</sub> are 9940 µg/L, 12740 µg/L by IV and PO respectively. The values of AUC are 9378, 25952 by IV and PO respectively<sup>[1]</sup>.

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

### PROTOCOL

#### Animal

Mice<sup>[1]</sup>

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**Administration** <sup>[1]</sup>

CDK8-IN-1 is administrated to mice with 5 mg/kg, 25mg/mg by IV and PO respectively<sup>[1]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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**REFERENCES**

[1]. Han X, et al. Discovery of potent and selective CDK8 inhibitors through FBDD approach. Bioorg Med Chem Lett. 2017 Sep 15;27(18):4488-4492.

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

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