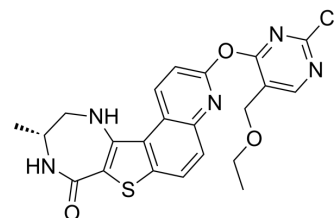


## Gamcemetinib

<b>Cat. No.:</b>	HY-139504		
<b>CAS No.:</b>	1887069-10-4		
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>20</sub> ClN <sub>5</sub> O <sub>3</sub> S		
<b>Molecular Weight:</b>	469.94		
<b>Target:</b>	MAPKAPK2 (MK2)		
<b>Pathway:</b>	MAPK/ERK Pathway		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 37.5 mg/mL (79.80 mM; ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	2.1279 mL	10.6397 mL	21.2793 mL
		5 mM	0.4256 mL	2.1279 mL	4.2559 mL
10 mM		0.2128 mL	1.0640 mL	2.1279 mL	
Please refer to the solubility information to select the appropriate solvent.					
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 1.5 mg/mL (3.19 mM); Suspended solution; Need ultrasonic				

### BIOLOGICAL ACTIVITY

<b>Description</b>	Gamcemetinib (CC-99677) is a potent, covalent, and irreversible inhibitor of the mitogen-activated protein (MAP) kinase-activated protein kinase-2 (MK2) pathway in both biochemical (IC <sub>50</sub> =156.3 nM) and cell based assays (EC <sub>50</sub> =89 nM). Gamcemetinib is extracted from patent WO2020236636, compound 1 <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	MK2 <sup>[1]</sup>
<b>In Vitro</b>	Gamcemetinib is a potent, covalent, and irreversible inhibitor of the mitogen-activated protein (MAP) kinase-activated protein kinase-2 (MK2) pathway in both biochemical (IC <sub>50</sub> =156.3 nM) and cell based assays (EC <sub>50</sub> =89 nM) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

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

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