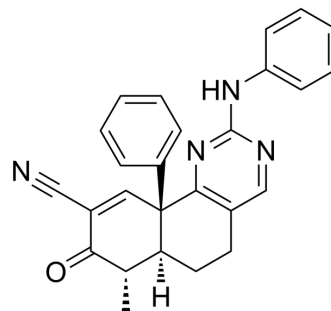


## IDH1 Inhibitor 2

<b>Cat. No.:</b>	HY-128661	
<b>CAS No.:</b>	2244895-42-7	
<b>Molecular Formula:</b>	C <sub>26</sub> H <sub>22</sub> N <sub>4</sub> O	
<b>Molecular Weight:</b>	406.48	
<b>Target:</b>	Isocitrate Dehydrogenase (IDH)	
<b>Pathway:</b>	Metabolic Enzyme/Protease	
<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 : 100 mg/mL (246.01 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	<b>Preparing Stock Solutions</b>	1 mM	2.4601 mL	12.3007 mL	24.6015 mL
		5 mM	0.4920 mL	2.4601 mL	4.9203 mL
10 mM		0.2460 mL	1.2301 mL	2.4601 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 >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (6.15 mM); Clear solution				

### BIOLOGICAL ACTIVITY

<b>Description</b>	IDH1 Inhibitor 2 (compound 13) is a potent wild-type IDH1 inhibitor via a direct covalent modification of His315, with an IC <sub>50</sub> of 110 nM <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	IDH1

### REFERENCES

[1]. Jakob CG, et al. Novel Modes of Inhibition of Wild-Type Isocitrate Dehydrogenase 1 (IDH1): Direct Covalent Modification of His315. J Med Chem. 2018 Aug 9;61(15):6647-6657.

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

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