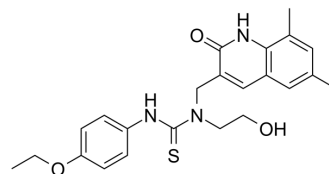


## β-Glucuronidase-IN-1

<b>Cat. No.:</b>	HY-103081		
<b>CAS No.:</b>	484006-66-8		
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>27</sub> N <sub>3</sub> O <sub>3</sub> S		
<b>Molecular Weight:</b>	425.54		
<b>Target:</b>	Bacterial		
<b>Pathway:</b>	Anti-infection		
<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 (235.00 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	2.3500 mL	11.7498 mL	23.4996 mL
		5 mM	0.4700 mL	2.3500 mL	4.6999 mL
10 mM		0.2350 mL	1.1750 mL	2.3500 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: ≥ 2.5 mg/mL (5.87 mM); Clear solution  2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (5.87 mM); Clear solution; Need ultrasonic				

### BIOLOGICAL ACTIVITY

<b>Description</b>	β-Glucuronidase-IN-1 is a potent, selective, uncompetitive, and orally active E. coli bacterial β-glucuronidase inhibitor, exhibiting an IC <sub>50</sub> and a K <sub>i</sub> of 283 nM and 164 nM, respectively <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	IC <sub>50</sub> : 283 nM (E. coli β-glucuronidase) K <sub>i</sub> : 164 nM (E. coli β-glucuronidase) <sup>[1]</sup>
<b>In Vitro</b>	β-Glucuronidase-IN-1 (Inhibitor 1) (0.01-100 μM) inhibits E. coli β-glucuronidase activity as a dose-dependent manner and exhibits an IC <sub>50</sub> and a K <sub>i</sub> value with 283 nM and 164 nM, respectively <sup>[1]</sup> . β-Glucuronidase-IN-1 (100 μM; 24-72 hours) maintains potent efficacy in living bacterial cells (EC <sub>50</sub> =17.7 nM), it does not affect bacterial cell growth under aerobic or anaerobic conditions or killing mammalian epithelial cells <sup>[1]</sup> .

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

#### Cell Viability Assay<sup>[1]</sup>

Cell Line:	CMT93, CaCO-2, HCT116 cells
Concentration:	100 $\mu$ M
Incubation Time:	24 hours, 48 hours and 72 hours
Result:	Did not impact mammalian cell survival and any reduction in cell viability were attributed to the presence of DMSO.

#### In Vivo

$\beta$ -Glucuronidase-IN-1 (oral gavage; 10  $\mu$ g; twice per day; 11 days) protects the mouse GI epithelium from CPT-11-induced damage and protects the glandular structure of CPT-11-treated intestinal tissues<sup>[1]</sup>.

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

Animal Model:	Healthy 6- to 8-week-old Balb/cJ mice <sup>[1]</sup>
Dosage:	10 $\mu$ g
Administration:	Oral gavage; twice per day; 11 days
Result:	Alleviated CPT-11-induced toxicity in mice.

## REFERENCES

[1]. Wallace BD, et al. Alleviating cancer drug toxicity by inhibiting a bacterial enzyme. *Science*. 2010 Nov 5;330(6005):831-5.

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

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