

## MM 419447

Cat. No.:	HY-P3282
CAS No.:	1092457-78-7
Molecular Formula:	C <sub>50</sub> H <sub>70</sub> N <sub>14</sub> O <sub>19</sub> S <sub>6</sub>
Molecular Weight:	1363.56
Sequence Shortening:	CCEYCCNPACTGC (Disulfide bridge:Cys1-Cys6; Cys2-Cys10; Cys5-Cys13)
Target:	Guanylate Cyclase
Pathway:	GPCR/G Protein
Storage:	Sealed storage, away from moisture
	Powder    -80°C    2 years
	-20°C    1 year

CCEYCCNPACTGC (Disulfide bridge:Cys1-Cys6; Cys2-Cys10; Cys5-Cys13)

\* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)

### SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (73.34 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
			1 mM	0.7334 mL	3.6669 mL	7.3337 mL
			5 mM	0.1467 mL	0.7334 mL	1.4667 mL
			10 mM	0.0733 mL	0.3667 mL	0.7334 mL
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (1.83 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (1.83 mM); Clear solution					

### BIOLOGICAL ACTIVITY

Description	MM 419447, a linaclotide metabolite, is a guanylate cyclase-C agonist. MM 419447 has the potential for the research of the irritable bowel syndrome with constipation (IBS-C) <sup>[1]</sup> .
In Vitro	MM 419447 exhibits high-affinity binding to T84 cells, resulting in a significant, concentration-dependent accumulation of intracellular cyclic guanosine-39,59-monophosphate (cGMP) <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	MM 419447 (10 mg/kg; p.o.) treatment shows the C <sub>max</sub> , AUC <sub>0-6</sub> and t <sub>1/2</sub> values of 27 ng/mL, ≤ 29.7 ng h/mL and 0.33 hours,

respectively<sup>[1]</sup>

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Animal Model:	Female Sprague-Dawley rats <sup>[1]</sup>
Dosage:	10 mg/kg
Administration:	P.o. (Pharmacokinetic Analysis)
Result:	The C <sub>max</sub> , AUC <sub>0-6</sub> and T <sub>max</sub> were 27 ng/mL, ≤ 29.7 ng h/mL and 0.33 hours, respectively.

## REFERENCES

[1]. Busby RW, et al. Pharmacologic properties, metabolism, and disposition of linaclotide, a novel therapeutic peptide approved for the treatment of irritable bowel syndrome with constipation and chronic idiopathic constipation. J Pharmacol Exp Ther. 2013;344

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

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