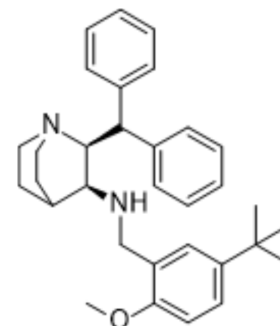


Maropitant

Cat. No.:	HY-10053		
CAS No.:	147116-67-4		
Molecular Formula:	C ₃₂ H ₄₀ N ₂ O		
Molecular Weight:	468.67		
Target:	Neurokinin Receptor		
Pathway:	GPCR/G Protein; Neuronal Signaling		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 25 mg/mL (53.34 mM); ultrasonic and warming and heat to 60°C)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	2.1337 mL	10.6685 mL	21.3370 mL
		5 mM	0.4267 mL	2.1337 mL	4.2674 mL
10 mM		0.2134 mL	1.0668 mL	2.1337 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	1. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (5.33 mM); Clear solution				

BIOLOGICAL ACTIVITY

Description	Maropitant is a selective and orally active neurokinin (NK1) receptor antagonist. Maropitant acts by blocking the binding of substance P within the emetic center and the chemoreceptor trigger zone (CRTZ). Maropitant is highly effective in preventing vomiting ^{[1][2]} .
IC ₅₀ & Target	NK1
In Vivo	Treatment with 1 mg/kg Maropitant citrate, significantly reduces the size of ulcerative dermatitis (UD) lesions in mice. Intravenous Maropitant decreases MAC by 16%. In contrast, epidural administration of either saline or Maropitant does not change the MAC (2.17% and 1.92 %, respectively). MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

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- [4]. Benchaoui HA, Cox SR, Schneider RP et al. The pharmacokinetics of maropitant, a novel neurokinin type-1 receptor antagonist, in dogs. *J Vet Pharmacol Ther.* 2007 Aug;30(4):336-44.
- [5]. Maropitant
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Caution: Product has not been fully validated for medical applications. For research use only.

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