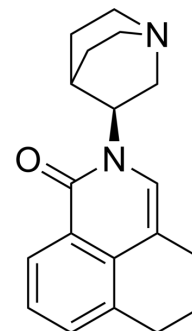


Dehydro Palonosetron

Cat. No.:	HY-44132
CAS No.:	135729-56-5
Molecular Formula:	C ₁₉ H ₂₂ N ₂ O
Molecular Weight:	294.39
Target:	5-HT Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Dehydro Palonosetron (RS 42358-197) is a potent, selective and orally active 5-HT ₃ receptor antagonist. Dehydro Palonosetron has no effect on the activities of 5-HT ₁ receptors, 5-HT ₂ receptors or 5-HT ₄ receptors ^[1] .								
IC₅₀ & Target	5-HT ₃ Receptor								
In Vitro	In functional experiments in vitro, Dehydro Palonosetron (RS 42358-197) behaves as a competitive antagonist against 5-HT-induced contractions in the guinea pig ileum, yielding a pA ₂ estimate of 8.1. Dehydro Palonosetron is devoid of any agonistic or antagonistic activity at 5-HT ₁ -like receptors, 5-HT ₂ receptors or 5-HT ₄ receptors ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
In Vivo	In anesthetized rats, Dehydro Palonosetron (RS 42358-197), administered by the intravenous, intraduodenal or transdermal route, dose-dependently inhibited the Bezold-Jarisch reflex induced by 2-methyl 5-HT (ID ₅₀ :0.05 µg/kg; i.v., 5.7 µg/kg; i.d., and 11.6 µg/chamber, respectively) ^[1] . Dehydro Palonosetron (RS 42358-197; 0.01 ng/kg-10 mg/kg; oral administration; once) disinhibits behaviour in the mouse suppressed by the aversive situation of the light/dark test box ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.								
	<table border="1"> <tr> <td>Animal Model:</td> <td>Male albino BKW mice (30-35 g)^[2]</td> </tr> <tr> <td>Dosage:</td> <td>0.01 ng/kg, 0.1 ng/kg, 1 ng/kg, 100 ng/kg, 1 µg/kg, 100 µg/kg, 1 mg/kg, 10 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral administration; once</td> </tr> <tr> <td>Result:</td> <td>Increased the proportion of time mice spent and the number of rearings and line crossings in the light area of the test chamber.</td> </tr> </table>	Animal Model:	Male albino BKW mice (30-35 g) ^[2]	Dosage:	0.01 ng/kg, 0.1 ng/kg, 1 ng/kg, 100 ng/kg, 1 µg/kg, 100 µg/kg, 1 mg/kg, 10 mg/kg	Administration:	Oral administration; once	Result:	Increased the proportion of time mice spent and the number of rearings and line crossings in the light area of the test chamber.
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Result:	Increased the proportion of time mice spent and the number of rearings and line crossings in the light area of the test chamber.								

REFERENCES

- [1]. R M Eglén, et al. RS 42358-197, a novel and potent 5-HT₃ receptor antagonist, in vitro and in vivo. *J Pharmacol Exp Ther.* 1993 Aug;266(2):535-43.
- [2]. B Costall, et al. The effect of the 5-HT₃ receptor antagonist, RS-42358-197, in animal models of anxiety. *Eur J Pharmacol.* 1993 Mar 30;234(1):91-9.

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

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