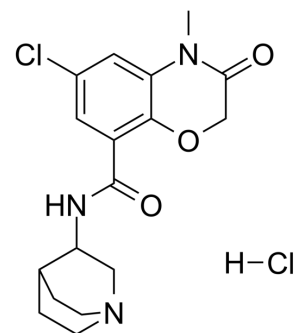


Azasetron hydrochloride

Cat. No.:	HY-B0068
CAS No.:	123040-16-4
Molecular Formula:	C ₁₇ H ₂₁ Cl ₂ N ₃ O ₃
Molecular Weight:	386.27
Target:	5-HT Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



SOLVENT & SOLUBILITY

In Vitro	H ₂ O : 20 mg/mL (51.78 mM; Need ultrasonic)																										
	DMSO : 2.22 mg/mL (5.75 mM; Need ultrasonic)																										
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent</th> <th rowspan="2">Mass</th> <th>1 mg</th> <th>5 mg</th> <th>10 mg</th> </tr> <tr> <th>Concentration</th> <th></th> <th></th> <th></th> </tr> </thead> <tbody> <tr> <td rowspan="4">Preparing Stock Solutions</td> <td>1 mM</td> <td>2.5889 mL</td> <td>12.9443 mL</td> <td>25.8886 mL</td> </tr> <tr> <td>5 mM</td> <td>0.5178 mL</td> <td>2.5889 mL</td> <td>5.1777 mL</td> </tr> <tr> <td>10 mM</td> <td>0.2589 mL</td> <td>1.2944 mL</td> <td>2.5889 mL</td> </tr> <tr> <td></td> <td></td> <td></td> <td></td> </tr> </tbody> </table>	Solvent	Mass	1 mg	5 mg	10 mg	Concentration				Preparing Stock Solutions	1 mM	2.5889 mL	12.9443 mL	25.8886 mL	5 mM	0.5178 mL	2.5889 mL	5.1777 mL	10 mM	0.2589 mL	1.2944 mL	2.5889 mL				
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Please refer to the solubility information to select the appropriate solvent.																											
In Vivo	1. Add each solvent one by one: PBS Solubility: 50 mg/mL (129.44 mM); Clear solution; Need ultrasonic																										

BIOLOGICAL ACTIVITY

Description	Azasetron (Y-25130) hydrochloride, a benzamide derivative, is a potent and selective 5-HT ₃ receptor antagonist. Azasetron is used in the study for Chemotherapy-induced nausea and vomiting (CINV) ^[1] .
IC₅₀ & Target	5-HT ₃ Receptor 0.33 nM (IC ₅₀)
In Vivo	Azasetron could effectively penetrate through the skin and pass into the systemic circulation ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
Animal Model:	Four male Bama miniature pigs weighing 9-11 kg (15-16 weeks old) ^[2] .
Dosage:	0.5 mg/kg.

Administration:	I.V. administration via the abdominal vein.
Result:	The mean plasma concentration dropped to minimum at 36 h. $C_{\max} = 44.88 \pm 7.16$ ng/mL) was achieved at the time point of 66.00 ± 22.98 h (T_{\max}).

CUSTOMER VALIDATION

- Protein Cell. 2019 Mar;10(3):178-195.

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REFERENCES

- [1]. Sato, N., et al., Antagonistic activity of Y-25130 on 5-HT₃ receptors. Jpn J Pharmacol, 1992. 59(4): p. 443-8.
- [2]. Haga, K., et al., The effects of orally administered Y-25130, a selective serotonin₃-receptor antagonist, on chemotherapeutic agent-induced emesis. Jpn J Pharmacol, 1993. 63(3): p. 377-83.

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

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