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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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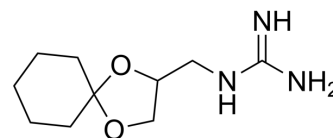
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Guanadrel

Cat. No.:	HY-W376701
CAS No.:	40580-59-4
Molecular Formula:	C ₁₀ H ₁₉ N ₃ O ₂
Molecular Weight:	213.28
Target:	Adrenergic Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (234.43 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		4.6887 mL	23.4434 mL	46.8867 mL
		5 mM		0.9377 mL	4.6887 mL	9.3773 mL
		10 mM		0.4689 mL	2.3443 mL	4.6887 mL
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (11.72 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (11.72 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil					
	Solubility: ≥ 2.5 mg/mL (11.72 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	Guanadrel is an orally active postganglionic adrenergic inhibitor of spiroketal. Guanadre can be used in anti-hypertensive studies ^[1] .	
In Vivo	Guanadrel (oral gavage, 15-400 mg/kg, every day, 6-52 weeks) has little toxicity to the nervous system in rats and dogs ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Rats, dogs ^[1]

Dosage:	15-400 mg/kg
Administration:	Oral gavage; every day; 6-52 weeks
Result:	Showed a slight increase in the weight of the prostate and adrenal glands while loss in the kidney in rats. Decreased weight gain values in a dose-dependent manner but liver function is normal in dogs.

REFERENCES

[1]. J D Palmer, et al. Guanadrel sulfate: a postganglionic sympathetic inhibitor for the treatment of mild to moderate hypertension. Pharmacotherapy. 1983 Jul-Aug;3(4):220-9. doi: 10.1002/j.1875-9114.1983.tb03257.x.

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

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