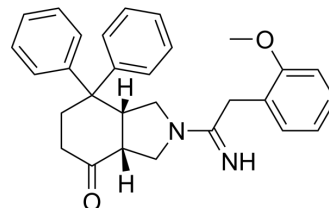


RP 67580

Cat. No.:	HY-108483
CAS No.:	135911-02-3
Molecular Formula:	C ₂₉ H ₃₀ N ₂ O ₂
Molecular Weight:	438.56
Target:	Neurokinin Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	RP 67580 is a non-peptide antagonist of substance P (SP), competitively inhibits the binding of [3H]SP to neurokinin receptor 1 (NK1 receptor) in rat brain membranes with a K _i value of 4.16 nM. RP 67580 is a specific antagonist of NK1 receptors and can be used in the research of pain and neurogenic inflammation ^[1] .								
In Vivo	<p>RP 67580 (0.03-1 mg/kg; i.v.; single dose) significantly inhibits neurogenic inflammation. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Male albino Sprague-Dawley rats</td> </tr> <tr> <td>Dosage:</td> <td>0.03, 0.1, 0.3, 1 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>i.v.; single dose</td> </tr> <tr> <td>Result:</td> <td>Potently and dose-dependently inhibited the strong plasma extravasation caused by electrical stimulation of the saphenous nerve for 15 minutes with an ED₅₀ value of 0.15 mg/kg.</td> </tr> </table>	Animal Model:	Male albino Sprague-Dawley rats	Dosage:	0.03, 0.1, 0.3, 1 mg/kg	Administration:	i.v.; single dose	Result:	Potently and dose-dependently inhibited the strong plasma extravasation caused by electrical stimulation of the saphenous nerve for 15 minutes with an ED ₅₀ value of 0.15 mg/kg.
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REFERENCES

[1]. Garret C, et al. Pharmacological properties of a potent and selective nonpeptide substance P antagonist. Proc Natl Acad Sci U S A. 1991 Nov 15;88(22):10208-12.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA