

Product Data Sheet

Product Name: Rispenzepine

Cat. No.: GC31035

Chemical Properties

Cas. No. 96449-05-7

SMILES O=C1C2=CC=CN=C2N(C(C3CN(C)CCC3)=O)C4=CC=CC=C4N1

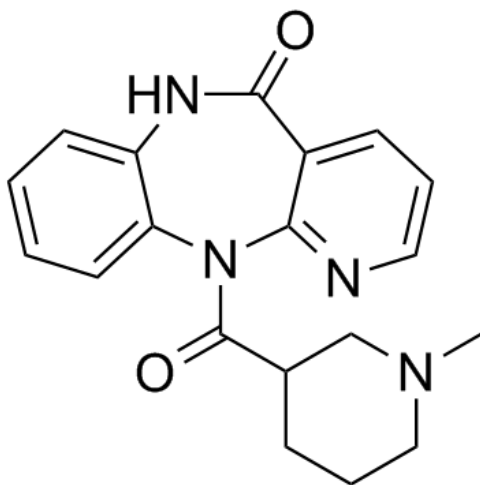
Formula $C_{19}H_{20}N_4O_2$ M.Wt 336.39

Solubility Soluble in DMSO Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure



Background

Rispenzepine is a novel antimuscarinic compound with a preferential action at M1, and M3 receptor subtypes.

The presence of muscarinic autoreceptors in human and guinea pig trachea is investigated by comparing the effects of the muscarinic receptor antagonists Pirenzepine (M1), Methoctramine (M2), 4-DAMP (M3), and Rispenzepine (M1/M3) on cholinergic neural contractile responses evoked by electrical field stimulation (EFS) and [3H]ACh release. The M1, M1/M3, or M3 antagonists inhibit the

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EFS-evoked cholinergic contractile response in a concentration-dependent manner (4-DAMP > Risperzepine > Pirenzepine), whereas Methoctramine facilitates this response at low concentrations (<3 μ M). In ACh release studies, the M3 antagonist has no significant effect, whereas Pirenzepine, Methoctramine, and Risperzepine significantly increase ACh release in guinea pig trachea. Risperzepine almost completely inhibits cholinergic, contractile responses at 0.3 μ M ($92.7 \pm 6.2\%$ inhibition, $n=6$, $p<0.05$; pD2 value of 7.31 ± 0.15) [1].

[1]. Patel HJ, et al. Evidence for prejunctional muscarinic autoreceptors in human and guinea pig trachea. Am J Respir Crit Care Med. 1995 Sep;152(3):872-8.

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