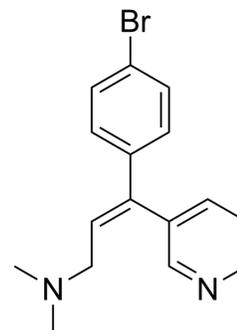


Zimelidine

Cat. No.:	HY-118835
CAS No.:	56775-88-3
Molecular Formula:	C ₁₆ H ₁₇ BrN ₂
Molecular Weight:	317.22
Target:	5-HT Receptor; Serotonin Transporter
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Zimelidine is a potent and selective inhibitor of serotonin 5-HT uptake and SERT. Zimelidine is an antidepressant ^{[1][2][3][4]} .																
In Vivo	<p>Zimelidine (15 mg/kg, IP, once) reduces the development of tolerance to morphine-induced antinociception in rats^[3]. Zimelidine (5 mg/kg, IP, daily for 14 days) dose not modify the responsiveness of CA₃ hippocampal pyramidal neurons to microiontophoretically applied 5-HT^[2].</p> <p>Zimelidine (0.2, 2 and 20 nmol/100 nL) in medial amygdaloid nucleus (MeA) evokes dose dependent hypophagic effects in fasted rats^[4].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tbody> <tr> <td>Animal Model:</td> <td>Male Wistar albino rats (160-180 g, n=72)^[3]</td> </tr> <tr> <td>Dosage:</td> <td>15 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>IP, once</td> </tr> <tr> <td>Result:</td> <td>Significantly attenuated the development and expression of morphine tolerance. The maximal antinociceptive effect of Zimelidine was obtained at the 60 minutes measurements in the zimelidine group and at the 30 minutes measurements in the morphine tolerant group by the tail-flick and hot-plate tests. Administration of zimelidine with morphine showed additive analgesic effect.</td> </tr> </tbody> </table> <table border="1"> <tbody> <tr> <td>Animal Model:</td> <td>Sprague-Dawley rats (150 to 250 g, Ten, male)^[2]</td> </tr> <tr> <td>Dosage:</td> <td>5 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>IP, daily for 14 days</td> </tr> <tr> <td>Result:</td> <td>Did not modify the responsiveness of CA₃ hippocampal pyramidal neurons to microiontophoretically applied 5-HT.</td> </tr> </tbody> </table>	Animal Model:	Male Wistar albino rats (160-180 g, n=72) ^[3]	Dosage:	15 mg/kg	Administration:	IP, once	Result:	Significantly attenuated the development and expression of morphine tolerance. The maximal antinociceptive effect of Zimelidine was obtained at the 60 minutes measurements in the zimelidine group and at the 30 minutes measurements in the morphine tolerant group by the tail-flick and hot-plate tests. Administration of zimelidine with morphine showed additive analgesic effect.	Animal Model:	Sprague-Dawley rats (150 to 250 g, Ten, male) ^[2]	Dosage:	5 mg/kg	Administration:	IP, daily for 14 days	Result:	Did not modify the responsiveness of CA ₃ hippocampal pyramidal neurons to microiontophoretically applied 5-HT.
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REFERENCES

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- [1]. Heel RC, et al. Zimelidine: a review of its pharmacological properties and therapeutic efficacy in depressive illness. *Drugs*. 1982 Sep;24(3):169-206.
- [2]. Blier P, et al. Electrophysiological investigations on the effect of repeated zimelidine administration on serotonergic neurotransmission in the rat. *J Neurosci*. 1983 Jun;3(6):1270-8.
- [3]. Ozdemir E, et al. Zimelidine attenuates the development of tolerance to morphine-induced antinociception. *Indian J Pharmacol*. 2012 Mar;44(2):215-8.
- [4]. Scopinho AA, et al. Medial amygdaloid nucleus 5-HT_{2c} receptors are involved in the hypophagic effect caused by zimelidine in rats. *Neuropharmacology*. 2012 Aug;63(2):301-9.
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Caution: Product has not been fully validated for medical applications. For research use only.

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