Product Data Sheet

Tipepidine

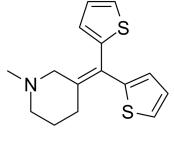
Cat. No.: HY-121685 CAS No.: 5169-78-8 Molecular Formula: C₁₅H₁₇NS₂ Molecular Weight: 275.43

Target: Potassium Channel

Pathway: Membrane Transporter/Ion Channel

Please store the product under the recommended conditions in the Certificate of Storage:

Analysis.



SOLVENT & SOLUBILITY

In Vitro

DMSO: 250 mg/mL (907.67 mM; Need ultrasonic)

	Solvent Mass Concentration	1 mg	5 mg	10 mg
reparing tock Solutions	1 mM	3.6307 mL	18.1534 mL	36.3069 mL
otock octations	5 mM	0.7261 mL	3.6307 mL	7.2614 mL
	10 mM	0.3631 mL	1.8153 mL	3.6307 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Tipepidine reversibly inhibits dopamine (DA) D_2 receptor-mediated GIRK currents ($I_{DA(GIRK)}$) with an IC_{50} of 7.0 μ M. Description

Tipepidine subsequently activates VTA dopamine neuron^[1]. Tipepidine, a non-narcotic antitussive, exerts an

antidepressant-like effect^[2].

IC50: 7.0 μM (dopamine D₂ receptor)^[1] IC₅₀ & Target

In Vivo Tipepidine (i.p.; 10-40 mg/kg; 0.5-23 hours) significantly decreases the immobility time in the forced swimming test in ACTH-

treated rats. Tipepidine (i.p.; 40 mg/kg) increases the extracellular dopamine level of the nucleus accumbens (NAc) in ACTHtreated rats^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model: Male Wistar rats weighting 150-240 g (5-7 weeks old) [2] Dosage: 10, 20 and 40 mg/kg. Administration: I.p.; 0.5, 5, 23 hours.

Result:	Decreased the immobility time in the forced swimming test in ACTH-treated rats.

REFERENCES

[1]. Hamasaki R, et al. Tipepidine activates VTA dopamine neuron via inhibiting dopamine D₂ receptor-mediated inward rectifying KM current. Neuroscience. 2013 Nov 12;252:24-34.

[2]. Kawaura K, et al. Tipepidine, a non-narcotic antitussive, exerts an antidepressant-like effect in the forced swimming test in adrenocorticotropic hormone-treated rats. Behav Brain Res. 2016 Apr 1;302:269-78.

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

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