Ispronicline

Cat. No.: HY-10063 CAS No.: 252870-53-4 Molecular Formula: $C_{14}H_{22}N_{2}O$ Molecular Weight: 234.34 Target: nAChR

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Pure form -20°C Storage: 3 years

2 years -80°C 6 months

In solvent -20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (426.73 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	4.2673 mL	21.3365 mL	42.6730 mL
	5 mM	0.8535 mL	4.2673 mL	8.5346 mL
	10 mM	0.4267 mL	2.1337 mL	4.2673 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 (10.67 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (10.67 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (10.67 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Ispronicline (TC-1734), an orally active, brain-selective α4β2 nicotine acetylcholine receptor (nAChR) partial agonist, has shown memory-enhancing properties in rodents and a good tolerability profile. Is pronicline binds to the $\alpha 4\beta 2$ nAChR with high affinity (K_i =11 nM) and is highly selective to other nAChRs such as α 7 nAChR and α 3 β 4 nAChR^{[1][2]}.

In Vivo

Ispronicline (10-20 mg/kg; s.c.) induces c-Fos expression in selective regions of the rat forebrain^[1]. Ispronicline pharmacokinetics (half-life of 2 h in rats) contrasts with the long-lasting improvement of working memory (18 h to 2 d)^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Wistar rats weighting 180-200 g ^[1]
Dosage:	10 and 30 mg/kg
Administration:	s.c.
Result:	Induced c-Fos in medial paraventricular nucleus of hypothalamus. Treatment with 30 mg/kg ispronicline, but not 10 mg/kg, increased the number of c-Fos immunoreactive cells significantly as compared to controls within the medial parvocellular part of paraventricular nucleus (PVN).

REFERENCES

[1]. Julie Jacobsen, et al. The α 4 β 2 nicotine acetylcholine receptor agonist is pronicline induces c-Fos expression in selective regions of the rat forebrain. Neurosci Lett. 2012 Apr 25;515(1):7-11.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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