

# Scopolamine (hydrobromide)

## **Catalog No: tcsc2000**

Available Sizes

**Size:** 100mg

Size: 500mg

Specifications

CAS No:

114-49-8

## Formula:

 $\mathsf{C}_{17}\mathsf{H}_{22}\mathsf{BrNO}_4$ 

**Pathway:** Neuronal Signaling;GPCR/G Protein;Neuronal Signaling;GPCR/G Protein

## **Target:**

mAChR;mAChR;5-HT Receptor;5-HT Receptor

#### **Purity / Grade:**

>98%

## Solubility: DMSO : $\geq$ 32 mg/mL (83.28 mM)

#### **Alternative Names:**

(-)-Scopolamine hydrobromide;Hyoscine hydrobromide;Scopine hydrobromide

## **Observed Molecular Weight:**

384.26

## **Product Description**

Scopolamine hydrobromide is a high affinity (nM) **muscarinic** antagonist. **5-HT**<sub>3</sub> receptor-responses are reversibly inhibited by Scopolamine with an IC<sub>50</sub> of 2.09  $\mu$ M.



IC50 & Target: IC50: 2.09 µM (5-HT<sub>3</sub> receptor)<sup>[1]</sup>

## mAChR<sup>[1]</sup>

*In Vitro:* Application of Scopolamine to oocytes expressing 5-HT<sub>3</sub> receptors does not elicit a response when applied alone, but causes a concentration-dependent inhibition of the response during a co-application of 2  $\mu$ M 5-HT. The plC<sub>50</sub> value for Scopolamine is 5.68±0.05 (IC<sub>50</sub>=2.09  $\mu$ M, n=6) with a Hill Slope of 1.06 ± 0.05. This gave a K<sub>b</sub> of 3.23  $\mu$ M. The same concentration-dependent effect is also seen when Scopolamine is applied during the 5-HT application. To further test for a competitive binding at the 5-HT<sub>3</sub> receptor, the competition of unlabelled Scopolamine is measured with [<sup>3</sup>H]granisetron, an established high-affinity competitive antagonist at these receptors. Scopolamine displays concentration-dependent competition with 0.6 nM [<sup>3</sup>H]granisetron (~K<sub>d</sub>), yielding an average pK<sub>i</sub> of 5.17±0.24 (K<sub>i</sub>=6.76  $\mu$ M, n=3)<sup>[1]</sup>.

In Vivo: In the histopathology study, there is no significant change in the histology of the brain. However, it is observed that there is a reduction in density of cells in the hippocampus of the control mice pretreated with Scopolamine who received only distilled water [2]. Scopolamine administration alone significantly increases the activity of Acetylcholinesterase enzyme (AchE) (7.98±0.065; P1-42) (P[3].



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