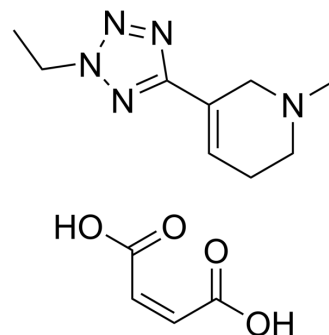


Alvamine maleate

| | |
|--------------------|---|
| Cat. No.: | HY-101586A |
| CAS No.: | 219581-36-9 |
| Molecular Formula: | C ₁₃ H ₁₉ N ₅ O ₄ |
| Molecular Weight: | 309.32 |
| Target: | mAChR |
| Pathway: | GPCR/G Protein; Neuronal Signaling |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

Description

Alvamine (Lu 25-109) maleate is a partial agonist of M1 and an antagonist of M2/M3. Alvamine maleate competitively antagonizes contractions induced by ammonium chloride (HY-Y1269) and electrical field stimulation in human detrusor muscle, indicating its potential application in regulating bladder function. Additionally, alvamine maleate can improve cognitive function following traumatic brain injury in rats^[1].

REFERENCES

- [1]. Jensen KG, et al. In vitro metabolism of the M1-muscarinic agonist 5-(2-ethyl-2H-tetrazol-5-yl)-1-methyl-1,2,3,6-tetrahydropyridine by human hepatic cytochromes P-450 determined at pH 7.4 and 8.5. *Drug Metab Dispos.* 1999 Jan;27(1):125-32.
- [2]. Waldeck K, et al. Actions of the new antimuscarinic compound Lu 25-109 on isolated human and pig detrusor. *Neurourol Urodyn.* 2002;21(1):92-8.
- [3]. Pike BR, et al. Chronic administration of a partial muscarinic M1 receptor agonist attenuates decreases in forebrain choline acetyltransferase immunoreactivity following experimental brain trauma. *Exp Neurol.* 1997 Sep;147(1):55-65

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

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