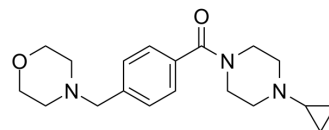


Bavisant

| | | | |
|---------------------------|---|-------|----------|
| Cat. No.: | HY-14880 | | |
| CAS No.: | 929622-08-2 | | |
| Molecular Formula: | C ₁₉ H ₂₇ N ₃ O ₂ | | |
| Molecular Weight: | 329.44 | | |
| Target: | Histamine Receptor | | |
| Pathway: | GPCR/G Protein; Immunology/Inflammation; Neuronal Signaling | | |
| Storage: | Powder | -20°C | 3 years |
| | | 4°C | 2 years |
| | In solvent | -80°C | 6 months |
| | | -20°C | 1 month |



SOLVENT & SOLUBILITY

| | | | | | |
|---|---|--------------------------|--------------|------------|------------|
| In Vitro | DMSO : 100 mg/mL (303.55 mM; Need ultrasonic) | | | | |
| | | Solvent Concentration | Mass 1 mg | 5 mg | 10 mg |
| | Preparing Stock Solutions | 1 mM | 3.0355 mL | 15.1773 mL | 30.3545 mL |
| | | 5 mM | 0.6071 mL | 3.0355 mL | 6.0709 mL |
| 10 mM | | 0.3035 mL | 1.5177 mL | 3.0355 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 (7.59 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (7.59 mM); Clear solution | | | | |

BIOLOGICAL ACTIVITY

| | |
|-------------------------------------|---|
| Description | Bavisant (JNJ-31001074) is an orally active, potent, brain-penetrating and highly selective antagonist of the histamine H ₃ receptor. Bavisant can be used for attention-deficit hyperactivity disorder (ADHD) research ^{[1][2][3]} . |
| IC₅₀ & Target | H ₃ receptor |
| In Vivo | Bavisant increases acetylcholine levels in rat frontal cortex ^[4] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

- [1]. Ghoshal A, et al. Identification of novel β -lactams and pyrrolidinone derivatives as selective Histamine-3 receptor (H3R) modulators as possible anti-obesity agents. *Eur J Med Chem.* 2018 May 25;152:148-159.
- [2]. Ghamari N, et al. Histamine H3 receptor antagonists/inverse agonists: Where do they go? *Pharmacol Ther.* 2019 Aug;200:69-84.
- [3]. Hudkins RL, et al. Discovery and characterization of 6-{4-[3-(R)-2-methylpyrrolidin-1-yl]propoxy}phenyl]-2H-pyridazin-3-one (CEP-26401, irdabisant): a potent, selective histamine H3 receptor inverse agonist. *J Med Chem.* 2011 Jul 14;54(13):4781-92.
- [4]. Weisler RH, et al. Randomized clinical study of a histamine H3 receptor antagonist for the treatment of adults with attention-deficit hyperactivity disorder. *CNS Drugs.* 2012 May 1;26(5):421-34.
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Caution: Product has not been fully validated for medical applications. For research use only.

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