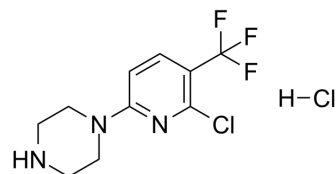


Org-12962 hydrochloride

| | |
|---------------------------|---|
| Cat. No.: | HY-21994 |
| CAS No.: | 210821-63-9 |
| Molecular Formula: | C ₁₀ H ₁₂ Cl ₂ F ₃ N ₃ |
| Molecular Weight: | 302.12 |
| Target: | 5-HT Receptor |
| Pathway: | GPCR/G Protein; Neuronal Signaling |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| | | | | | | | | | | | |
|-------------------------------------|---|---|---|---------------|---------------------------------|---------|-----------------------------------|-----------------|--|---------|---|
| Description | Org 12962 hydrochloride is a potent, selective and efficacious 5-HT _{2C} receptor agonist and exhibits pEC ₅₀ values of 7.01, 6.38 and 6.28 for 5-HT _{2C} , 5-HT _{2A} and 5-HT _{2B} , respectively. Org 12962 hydrochloride is effective in panic-like anxiety animal model [2]. | | | | | | | | | | |
| IC₅₀ & Target | 5-HT _{2C} Receptor 7.01 (pEC50) | 5-HT _{2A} Receptor 6.38 (pEC50) | 5-HT _{2B} Receptor 6.28 (pEC50) | | | | | | | | |
| In Vivo | <p>Org 12962 (intraperitoneal injection; 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg) displays antiaversive effects in a rat model of panic-like anxiety, it increases the postinjection frequency thresholds for self-interruption (F_{3,71}=11.40). The dose of 0.3 mg/kg ip is not significantly active, that the dose of 1 mg/kg is marginally active and that 3.2 mg/kg ip induced highly significant antiaversive effects^[3].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Male Wistar rats^[3]</td> </tr> <tr> <td>Dosage:</td> <td>0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intraperitoneal injection; 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg</td> </tr> <tr> <td>Result:</td> <td>Had a dose-related antipanic-like effect in SD-rat.</td> </tr> </table> | | | Animal Model: | Male Wistar rats ^[3] | Dosage: | 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg | Administration: | Intraperitoneal injection; 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg | Result: | Had a dose-related antipanic-like effect in SD-rat. |
| Animal Model: | Male Wistar rats ^[3] | | | | | | | | | | |
| Dosage: | 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg | | | | | | | | | | |
| Administration: | Intraperitoneal injection; 0.3 mg/kg, 1 mg/kg, and 3.2 mg/kg | | | | | | | | | | |
| Result: | Had a dose-related antipanic-like effect in SD-rat. | | | | | | | | | | |

REFERENCES

- [1]. Porter RH, et al. Functional characterization of agonists at recombinant human 5-HT_{2A}, 5-HT_{2B} and 5-HT_{2C} receptors in CHO-K1 cells. *Br J Pharmacol*. 1999 Sep;128(1):13-20.
- [2]. Jenck F, et al. Antiaversive effects of 5HT_{2C} receptor agonists and fluoxetine in a model of panic-like anxiety in rats. *Eur Neuropsychopharmacol*. 1998 Aug;8(3):161-8.
- [3]. Faassen F, et al. Caco-2 permeability, P-glycoprotein transport ratios and brain penetration of heterocyclic drugs. *Int J Pharm*. 2003 Sep 16;263(1-2):113-22.

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

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