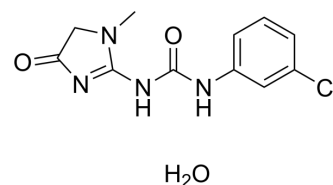


Fenobam hydrate

Cat. No.:	HY-101478A
CAS No.:	63540-28-3
Molecular Formula:	C ₁₁ H ₁₃ ClN ₄ O ₃
Molecular Weight:	284.7
Target:	mGluR; Apoptosis
Pathway:	GPCR/G Protein; Neuronal Signaling; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Fenobam hydrate is a selective and orally active mGluR5 antagonist (IC ₅₀ =84 nM) that can penetrate the blood-brain barrier. Fenobam hydrate shows the K _d values of 54 nM and 31 nM on rat and human recombinant mGlu5 receptors, respectively. Fenobam hydrate has anxiolytic activity, inhibits self-administration behavior in rat, and induces apoptosis in cancer cells. Fenobam hydrate can be used for research on neurological diseases, cancer and drug addiction ^{[1][2][3]} .										
IC₅₀ & Target	mGluR5 84 nM (IC ₅₀)	human mGluR5 31 nM (K _d)	rat mGluR5 54 nM (K _d)								
In Vitro	<p>Fenobam hydrate (300 μM; 72 h) significantly inhibits proliferation and induces apoptosis in LM7 cells^[2]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Apoptosis Analysis^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>LM7 cells</td> </tr> <tr> <td>Concentration:</td> <td>300 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Significantly reduced total number of cells, proliferating cells, and induced apoptosis.</td> </tr> </table>			Cell Line:	LM7 cells	Concentration:	300 μM	Incubation Time:	72 h	Result:	Significantly reduced total number of cells, proliferating cells, and induced apoptosis.
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Concentration:	300 μM										
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Result:	Significantly reduced total number of cells, proliferating cells, and induced apoptosis.										
In Vivo	<p>Fenobam hydrate (30-60 mg/kg; p.o.; 3 times a week) significantly inhibits self-administration behavior in rats^[3]. 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 Long-Evans rats (250-300 g)^[3].</td> </tr> <tr> <td>Dosage:</td> <td>30-60 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral administration; 3 times a week.</td> </tr> <tr> <td>Result:</td> <td>Inhibited self-administration.</td> </tr> </table>			Animal Model:	Male Long-Evans rats (250-300 g) ^[3] .	Dosage:	30-60 mg/kg	Administration:	Oral administration; 3 times a week.	Result:	Inhibited self-administration.
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REFERENCES

[1]. Porter RH, et al. Fenobam: a clinically validated nonbenzodiazepine anxiolytic is a potent, selective, and noncompetitive mGlu5 receptor antagonist with inverse agonist activity. *J Pharmacol Exp Ther.* 2005 Nov;315(2):711-21.

[2]. Liao S, et al. Osteosarcoma cell proliferation and survival requires mGluR5 receptor activity and is blocked by Riluzole. *PLoS One.* 2017 Feb 23;12(2):e0171256.

[3]. Keck TM, et al. Fenobam sulfate inhibits cocaine-taking and cocaine-seeking behavior in rats: implications for addiction treatment in humans. *Psychopharmacology (Berl).* 2013;229(2):253-265.

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

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