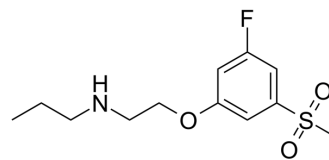


Mesdopetam

Cat. No.:	HY-109150
CAS No.:	1403894-72-3
Molecular Formula:	C ₁₂ H ₁₈ FNO ₃ S
Molecular Weight:	275.34
Target:	Dopamine Receptor
Pathway:	GPCR/G Protein; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Mesdopetam (IRL790) is a dopamine D ₃ receptor antagonist (K _i =90 nM; IC ₅₀ =9.8 μM for human recombinant D ₃ receptor) with psychomotor stabilizing properties. Mesdopetam is used for the research of motor and psychiatric complications in Parkinson disease ^{[1][2]} .								
IC₅₀ & Target	D ₃ Receptor								
In Vivo	<p>Mesdopetam (IRL790) (3.7, 11, 33, or 100 μmol/kg) dose-dependently inhibits the behavioral activation following pretreatment with D-amphetamine or MK-80^[1].</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 Sprague-Dawley rats^[1]</td> </tr> <tr> <td>Dosage:</td> <td>3.7, 11, 33, or 100 μmol/kg (synthesized in-house as HCl salt, was dissolved in physiologic saline (0.9% w/v NaCl))</td> </tr> <tr> <td>Administration:</td> <td>s.c. was administered subcutaneously 4 min before the start of recording</td> </tr> <tr> <td>Result:</td> <td>Dose-dependently inhibited the behavioral activation following pretreatment with D-amphetamine or MK-801.</td> </tr> </table>	Animal Model:	Male Sprague-Dawley rats ^[1]	Dosage:	3.7, 11, 33, or 100 μmol/kg (synthesized in-house as HCl salt, was dissolved in physiologic saline (0.9% w/v NaCl))	Administration:	s.c. was administered subcutaneously 4 min before the start of recording	Result:	Dose-dependently inhibited the behavioral activation following pretreatment with D-amphetamine or MK-801.
Animal Model:	Male Sprague-Dawley rats ^[1]								
Dosage:	3.7, 11, 33, or 100 μmol/kg (synthesized in-house as HCl salt, was dissolved in physiologic saline (0.9% w/v NaCl))								
Administration:	s.c. was administered subcutaneously 4 min before the start of recording								
Result:	Dose-dependently inhibited the behavioral activation following pretreatment with D-amphetamine or MK-801.								

REFERENCES

[1]. Waters S, et al. Preclinical Pharmacology of [2-(3-Fluoro-5-Methanesulfonyl-phenoxy)Ethyl](Propyl)amine (IRL790), a Novel Dopamine Transmission Modulator for the Treatment of Motor and Psychiatric Complications in Parkinson Disease. *J Pharmacol Exp Ther.* 2020;374(1):113-125.

[2]. Becanovic K, et al. Effects of a Novel Psychomotor Stabilizer, IRL790, on Biochemical Measures of Synaptic Markers and Neurotransmission. *J Pharmacol Exp Ther.* 2020;374(1):126-133.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA