

Product Data Sheet

Idalopirdine

Cat. No.: HY-14338 CAS No.: 467459-31-0 Molecular Formula: $C_{20}H_{19}F_{5}N_{2}O$ Molecular Weight: 398.37

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 Idalopirdine (Lu AE58054) is a potent, selective 5-HT6 receptor antagonist with a K_i value of 0.83 nM. Idalopirdine may be

 $used \ in \ studies \ of \ Alzheimer's \ disease \ and \ schizophrenia, \ among \ other \ related \ disorders \ ^{[1][2]}.$

IC₅₀ & Target 5-HT₆ Receptor

0.83 nM (Ki)

In Vivo Idalopirdine (intraperitoneal injection, 5 mg/kg, daily, 28 days) can reduce food intake and body weight in over-eating rat

> Idalopirdine (1 or 2 mg/kg, i.v) can dose-dependently increase the gamma power during nPO electrical stimulation, enhance effect of donepezil on cortical gamma oscillations but no alteration of sleep-wake patterns in rats^[2].

Animal Model:	Male Wistar rats ^[1]
Dosage:	5 mg/kg
Administration:	intraperitoneal injection, daily, 28 days
Result:	Significantly reduced the amount of calories consumed by animals in a palatable diet and significantly reduced plasma levels of glucose, triglycerides and cholesterol.
Animal Model:	Male Sprague-Dawley rats ^[2]
Dosage:	1 or 2 mg/kg
Administration:	i.v.
Result:	No significant increase in the gamma power at 1 mg/kg and significantly increased the gamma power at 2 mg/kg. Significantly enhanced and/or prolonged effect of low-dose donepezil (0.3 mg/kg) on gamma power during 60-minute nPO stimulation after donepezil administration.

REFERENCES

- [1]. Magdalena Kotańska, et al. Idalopirdine, a selective 5-HT6 receptor antagonist, reduces food intake and body weight in a model of excessive eating. Metab Brain Dis. 2018 Jun;33(3):733-740.
- [2]. Maria Amat-Foraster, et al. The 5-HT6 receptor antagonist idalopirdine potentiates the effects of donepezil on gamma oscillations in the frontal cortex of anesthetized and awake rats without affecting sleep-wake architecture. Neuropharmacology. 2017 Feb;113(Pt A):45-5

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

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