Product Data Sheet

Progabide

Molecular Weight:

 Cat. No.:
 HY-A0173

 CAS No.:
 62666-20-0

 $\label{eq:molecular Formula: C17H16ClFN2O2} \textbf{Molecular Formula:} \qquad \textbf{C}_{17}\textbf{H}_{16}\textbf{ClFN}_2\textbf{O}_2$

Target: GABA Receptor

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Storage: Powder -20°C 3 years

334.77

4°C 2 years

In solvent -80°C 6 months

-20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 250 mg/mL (746.78 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9871 mL	14.9356 mL	29.8713 mL
	5 mM	0.5974 mL	2.9871 mL	5.9743 mL
	10 mM	0.2987 mL	1.4936 mL	2.9871 mL

Please refer to the solubility information to select the appropriate solvent.

BIOL	α CI	~ 1	ACTI	MTV
вил	10/61	LAI	$\Delta U = I$	$\mathbf{v} - \mathbf{v}$

Description	Progabide is a gamma-aminobutyric acid receptor (GABA) agonist.
IC ₅₀ & Target	GABA receptor ^[1]
In Vivo	Progabide is a gamma-aminobutyric acid receptor (GABA) agonist. Doses of 50, 100 and 200 mg/kg Progabide given IP to male Wistar rats increase the plasma corticosterone levels by 244, 365 and 476% respectively (t=6.44 to12.55, p<0.01). 10 mg/kg of Progabide fails to change the concentration of corticosterone in plasma (t=0.76, N.S.). The increased plasma corticosterone level induced by administration of 200 mg/kg Progabide is evident 30 (t=2.625, p<0.05), 60 (t=13.13, p<0.001) and 120 min (t=4.07, p<0.01) after drug injection, but returns to the control value 240 min after drug injection (t=0.86, N.S.). The greatest corticosterone rise (compare with the corresponding control) is reached 60 min following the administration of Progabide ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

Animal
Administration [1]

Male Wistar rats weighing 170 to 240 g are caged in groups of three under diurnal lighting conditions with free access to food and water. The rats are accustomed to handling (housing 1 per cage for 30 min, injecting with saline, 1 mL/100 g body weight) in the period of 7 days before the beginning of the experiment. Progabide is injected as a suspension in saline containing 0.1% Tween 80. Control rats are treated IP with the corresponding vehicle (1 mL/100 g body weight). Ether stress is performed and 15 min after the beginning of the stressful procedure the animals are sacrificed^[1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Manev H, et al. Progabide, a GABA mimetic drug, stimulates the secretion of plasma corticosterone in rats. Pharmacol Biochem Behav. 1987 Dec;28(4):443-6.

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

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