

Neu2000

Catalog No: tcsc0025740



Available Sizes

Size: 1mg

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg



Specifications

CAS No:

640290-67-1

Formula:

$C_{15}H_8F_7NO_3$

Pathway:

Membrane Transporter/Ion Channel;Neuronal Signaling

Target:

iGluR;iGluR

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 112.5 mg/mL (293.57 mM)

Observed Molecular Weight:

383.22

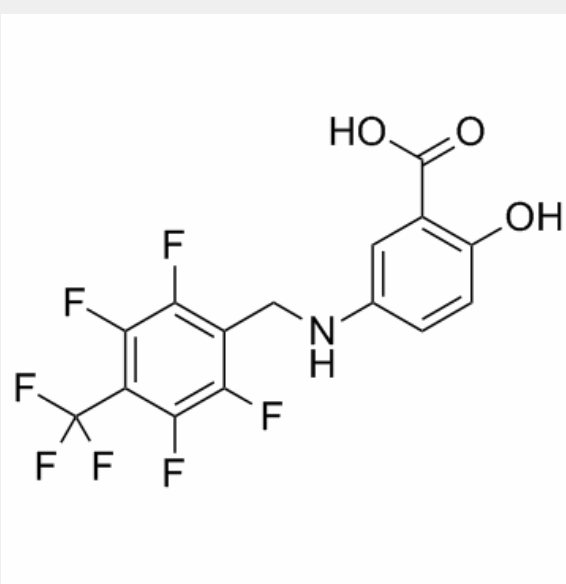
Product Description

Neu2000 is an uncompetitive **N-methyl-D-aspartate (NMDA)** receptor antagonist.

IC₅₀ & Target: NMDA receptor^[1]

In Vitro: Neu2000 shows apparent neuroprotection against 300 μ M N-methyl-D-aspartate (NMDA) at doses as low as 30 μ M. Neu2000 does not protect cortical neurons against α -amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid- or kainate-mediated excitotoxicity. Neu2000 inhibits the electrophysiologic response of cultured cortical neurons to 300 μ M NMDA in a concentration-dependent manner, indicating that the effect is mediated by a specific action at NMDA receptors. The Neu2000 dose-response has an IC₅₀ of 35.38 ± 5.94 μ M and Hill's coefficient of 0.91 (n=8). Neu2000 (100 μ M) significantly reduces the maximal NMDA response by $58.31 \pm 2.72\%$ (n=5) and the EC₅₀ values of NMDA from 18.88 ± 1.85 to 9.92 ± 0.17 μ M (n=5, P[1]).

In Vivo: Pharmacokinetic analysis reveals that the half-life of Neu2000 is 1.42, 2.14, and 1.79 h following intraperitoneal administration of 10, 25, and 50 mg/kg, respectively. In addition, the C_{max} (maximum plasma concentration) is calculated as 3.86, 18.73, and 52.83 μ g/mL and the AUC (area under the curve) is determined to be 7.37, 55.15, and 96.77 μ g/h/mL at the same respective doses. The levels of basal mitochondrial ROS are significantly elevated at 24 h post-surgery in both the vehicle-treated (4.1-fold, p[2]).



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