



Omberacetam

Catalog No: tcsc1575

Available Sizes
Size: 5mg
Size: 10mg
Size: 25mg
Size: 50mg
Size: 100mg
Specifications
CAS No: 157115-85-0
Formula: $C_{17}^{H}_{22}^{N}_{2}^{O}_{4}$
Pathway: Membrane Transporter/Ion Channel;Neuronal Signaling
Target: iGluR;iGluR
Purity / Grade: >98%
Solubility: DMSO : ≥ 100 mg/mL (314.10 mM)
Alternative Names:





Observed Molecular Weight:

318.37

Product Description

Omberacetam (GVS-111) is a medication promoted and prescribed in Russia and neighbouring countries as a nootropic.

In Vitro: Nooglutil exhibits pharmacologically significant competition with a selective agonist of AMPA receptors ([G-3H]Ro 48-8587) for the receptor binding sites (with IC50 = 6.4 + /-0.2 microM), while the competition of noopept for these receptor binding sites was lower by an order of magnitude (IC50 = 80 + /-5.6 microM) [1]. GVS-111 significantly increased neuronal survival after H(2)O(2)-treatment displaying a dose-dependent neuroprotective activity from 10 nM to 100 microM, and an IC(50) value of 1.21 + /-0.07 microM. GVS-111 inhibited the accumulation of intracellular free radicals and lipid peroxidation damage in neurons treated with H(2)O(2) or FeSO(4), suggesting an antioxidant mechanism of action [2].

In Vivo: N-Phenylacetyl-L-prolylglycine ethyl ester (GVS-111) administered intravenously at a dose of 0.5 mg/kg/day, for the first time 1 h after ischaemic lesion and then for 9 post-operative days, with the last administration 15 min before testing, attenuated the deficit [3]. GVS-111 itself was not found in rat brain 1 h after 5 mg/kg i.p. administration up to limit of detection (LOD) under high performance liquid chromatography (HPLC) conditions [4]. The most pronounced antiinflammatory effect of dipeptide was observed on the model of adjuvant arthritis in rats, where the drug administered over 25 days in a daily dose of 0.5 mg/kg (i.m.) or 5 mg/kg (p.o.) significantly reduced the chronic immune inflammation (on the 12th day, by 94.0 and 74.1%, respectively) [5].

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