

Product Data Sheet

CNS 5161

Cat. No.:HY-101809ACAS No.:160754-76-7Molecular Formula: $C_{16}H_{18}CIN_3S_2$ Molecular Weight:351.92

Target: iGluR

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	CNS-5161 is a novel NMDA ion-channel antagonist that interacts with the NMDA receptor/ion channel site to produce a noncompetitive blockade of the actions of glutamate.
IC ₅₀ & Target	$NMDA\ receptor/ion\ channel^{[1]}$
In Vitro	CNS-5161 (CNS 5161) is a novel and selective noncompetitive antagonist of the NMDA subset of glutamate receptors in the mammalian brain. CNS-5161 has potent inhibitory activity in vitro at the NMDA ion channel and is able to displace [³ H] MK-801 binding with a K _i of 1.8 nM in synaptosomal membrane preparations from rat brain ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	In the neonatal rat NMDA excitotoxicity model in vivo, CNS-5161 (CNS 5161) protects against the necrotic effects of exogenous N-methyl-D-aspartate with an ED ₈₀ of 4 mg/kg by the intraperitoneal (i.p.) route. CNS-5161 also shows a 91% inhibition of audiogenic seizures in DBA/2 mice at 4 mg/kg i.p., and has a neuroprotective effect following hypoxix/ischaemic brain injury in neonatal rats ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Walters MR, et al. Early clinical experience with the novel NMDA receptor antagonist CNS 5161. Br J Clin Pharmacol. 2002 Mar; 53(3):305-11.

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

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