Product Data Sheet

CP-465022 maleate

Cat. No.: HY-18663A
CAS No.: 199656-46-7
Molecular Formula: $C_{30}H_{28}ClFN_4O_5$

Molecular Weight: 579.02
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	CP-465022 Maleate is a potent, and selective noncompetitive AMPA receptor antagonist with anticonvulsant activity. CP-465022 is against Kainate-induced response with an IC ₅₀ of 25 nM in rat cortical neurons. CP-465022 provides a new tool to investigate the role of AMPA receptors in physiological and pathophysiological processes ^{[1][2]} .
IC ₅₀ & Target	IC50: 25 nM (rat cortical neurons) ^[1]
In Vitro	CP-465022 (0.0001 μ M-10 μ M) inhibits kainate-induced response in relatively slow manner and dependents on compound concentration, exhibiting a calculated IC ₅₀ of 25 nM and essentially complete inhibition at 3.2 μ M ^[1] . CP-465022 1 μ M for 10 min has little effect on peak NMDA-induced currents but reduces current measured at 8 s during NMDA application by 26%.CP-465,022 at 10 μ M inhibits peak NMDA-induced currents in cortical neurons by 36% and currents measured at 8 s by 70% d in primary cultures of cortical and cerebellar granule neurons ^[1] . CP-465022 1 μ M for 10 min inhibits peak NMDA currents in cultured rat cerebellar granule neurons with mean inhibition of 19% and NMDA currents measured at 8 s by 45%, similar to what is observed in the cortical neurons ^[1] . CP-465022 (100 nM -10 μ M) has inhibitory effects on Kainate-induced whole-cell currents in voltage-clamped rat hippocampal, 100 nM CP465,022 inhibits kainate currents developed over the course of 200s, 500 nM and 1 μ M CP-465,022 nearly complete inhibits this time frame (99.3%) ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. JT Lazzaro, et al. Functional characterization of CP-465,022, a selective, noncompetitive AMPA receptor antagonist. Neuropharmacology. 2002 Feb;42(2):143-53.

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

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