Product Data Sheet

NS3763

 $\begin{array}{lll} \textbf{Cat. No.:} & \text{HY-107603} \\ \textbf{CAS No.:} & 70553-45-6 \\ \textbf{Molecular Formula:} & \textbf{C}_{22}\textbf{H}_{16}\textbf{N}_2\textbf{O}_6 \\ \end{array}$

Molecular Weight: 404.37
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	NS3763 is a selective and noncompetitive GLU_{K5} receptor antagonist with an IC_{50} of 1.6 μ M. NS3763 does not show significant antagonistic properties on GLU_{K6} , AMPA or NMDA receptors ^[1] .
IC ₅₀ & Target	IC50: 1.6 μ M (GLU _{K5} receptor) ^[1]
In Vitro	In human embryonic kidney HEK293 cells expressing homomeric GLU_{K5} or GLU_{K6} receptors, NS3763 displays selectivity for inhibition of domoate-induced increase in intracellular calcium mediated through the $GLUK5$ subtype ($IC_{50} = 1.6 \mu M$) of kainate receptors compared with the $GLUK6$ subtype ($IC_{50} > 30 \mu M$) ^[1] . NS3763 selectively inhibits l-glutamate- and domoate-evoked currents through GLU_{K5} receptors in HEK293 cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Jeppe K Christensen, et al. In vitro characterization of 5-carboxyl-2,4-di-benzamidobenzoic acid (NS3763), a noncompetitive antagonist of GLUK5 receptors. J Pharmacol Exp Ther. 2004 Jun;309(3):1003-10.

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

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