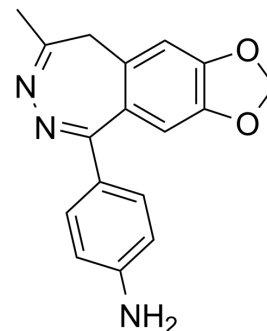


GYKI 52466

Cat. No.:	HY-103234
CAS No.:	102771-26-6
Molecular Formula:	C ₁₇ H ₁₅ N ₃ O ₂
Molecular Weight:	293.32
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	GYKI 52466 is an orally active, highly selective and noncompetitive AMPA/kainate receptor antagonist with the IC ₅₀ values of 7.5 and 11 μM, respectively. GYKI 52466 has good blood brain barrier permeability and anticonvulsant effect. GYKI 52466 can be used in Parkinson's disease research ^{[1][2]} .	
IC₅₀ & Target	AMPA Receptor 7.5 μM (IC ₅₀)	Kainate Receptor 11 μM (IC ₅₀)
In Vitro	GYKI 52466 (0.3-100 μM) inhibits inward currents activated by AMPA and Kainate receptor in cultured rat hippocampal neurons ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
In Vivo	GYKI 52466 (intraperitoneal injection; 1.76-13.2 mg/kg; once) treatment provides potent anticonvulsant protection against sound-induced seizures in DBA/2 mice ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Male and female DBA/2 mice tested for sound-induced seizure responses ^[2]
	Dosage:	1.76-13.2 mg/kg
	Administration:	Intraperitoneal injection; 1.76-13.2 mg/kg; once
	Result:	Observed Maximal anticonvulsant protection after the i.p. treatment (5-15 min).

REFERENCES

- [1]. S D Donevan, et al. GYKI 52466, a 2,3-benzodiazepine, is a highly selective, noncompetitive antagonist of AMPA/kainate receptor responses. *Neuron*. 1993 Jan;10(1):51-9.
- [2]. A G Chapman, et al. The anticonvulsant effect of the non-NMDA antagonists, NBQX and GYKI 52466, in mice. *Epilepsy Res*. 1991 Jul;9(2):92-6.

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

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