Proteins

Product Data Sheet

gamma-DGG

Target:

Cat. No.: HY-100785 CAS No.: 6729-55-1 Molecular Formula: $C_7 H_{12} N_2 O_5$ Molecular Weight: 204.18 iGluR

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

-20°C Storage: Powder 3 years

4°C 2 years

-80°C In solvent 2 years

> -20°C 1 year

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SOLVENT & SOLUBILITY

 $H_2O : \ge 150 \text{ mg/mL} (734.65 \text{ mM})$ In Vitro

* "≥" means soluble, but saturation unknown.

	Solvent Mass Concentration	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	4.8976 mL	24.4882 mL	48.9764 mL
	5 mM	0.9795 mL	4.8976 mL	9.7953 mL
	10 mM	0.4898 mL	2.4488 mL	4.8976 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

In Vitro

1. Add each solvent one by one: PBS

Solubility: 25 mg/mL (122.44 mM); Clear solution; Need ultrasonic and warming and heat to 60°C

BIOLOGICAL ACTIVITY

Description gamma-DGG is a competitive AMPA receptor blocker.

IC₅₀ & Target AMPA receptor^[1]

> gamma-DGG (γ -DGG), a competitive AMPA receptor blocker that blocks less at higher glutamate concentration. At 200-400 μ M, gamma-DGG in the bath reduces the miniature EPSC (mEPSC) amplitude by 26±2% (n=5 synapses), and shifts both the mEPSC amplitude distribution and the cumulative probability curve to the left [1], gamma-DGG (γ -DGG) is the most effective antagonist of the excitatory post-synaptic potentials (e.p.s.p.s). Its action is reversible and not associated with any change in the passive membrane properties of the granule cells or in the apparent reversal potential of the e.p.s.p. Quantal analysis shows that the reduction in the e.p.s.p. paralleled the decrease in quantal size rather than quantal content, confirming a post-synaptic site of the action of gamma-DGG^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Wu XS, et al. The origin of quantal size variation: vesicular glutamate concentration plays a significant role. J Neurosci. 2007 Mar 14;27(11):3046-56.

[2]. Crunelli V, et al. Blockade of amino acid-induced depolarizations and inhibition of excitatory post-synaptic potentials in rat dentate gyrus. J Physiol. 1983 Aug;341:627-40.

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

Tel: 609-228-6898 Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA

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