Hello Bio, Inc. 304 Wall St., Princeton, NJ 08540 USA

T. 609-683-7500 F. 609-228-4994

customercare-usa@hellobio.com



DATASHEET CNQX disodium salt

Product overview

Name	CNQX disodium salt
Cat No	HB0205
Biological action	Antagonist
Purity	>98%
Customer comments	The CNQX is going fine ! Verified customer, IBPS, Inserm, CNRS
	It works exactly as it should! Dissolved in water, kept in aliquots in -20 freezer. Verified customers,

Description

It works exactly as it should! Dissolved in water, kept in aliquots in -20 freezer. Verified custon SickKids (University of Toronto) Potent, competitive AMPA / kainate receptor antagonist. Disodium salt.

Images



Biological Data

Biological description	CNQX disodium salt is a water soluble, potent and competitive AMPA and kainate receptor antagonist. CNQX also antagonizes NMDA receptors at the glycine site.
	CNQX increases GABA _A receptor spontaneous postsynaptic currents (sPSCs) and also shows neuroprotective actions.
Application notes	$\frac{\text{CNQX}}{\text{CNQX}}$ also available. The AMPA receptor antagonist CNQX disodium salt is commonly used at concentrations of 10 μM to inhibit the actions of glutamate acting on AMPARs.
	CNQX disodium salt from Hello Bio reduces both spontaneous and evoked EPSCs in cortical neurons at concentrations of 1 μ M with full AMPA receptor blockade at 10 μ M (see Fig 1 above).

#Protocol 1: Evoked and spontaneous excitatory post synaptic currents (EPSCs)

- Whole cell voltage clamp recordings were obtained from layer V neurons of the mouse prelimbic cortex brain slice.
- EPSCs were evoked via a stimulating electrode placed in layers II/III delivering a single square (150 μ s) pulse every 10 sec at an intensity that gave a reliable EPSC.
- Neurons were held at -70 to -60 mV (the reversal potential of GABA currents).

EPSCs were continuously stimulated and recorded in response to 5 min applications of varying concentrations of CNQX disodium salt until complete receptor inhibition.

- Spontaneous EPSCs were recorded before and after addition of CNQX disodium salt by holding the neuron at -70 mV and recording for 10 sec.
- Recordings for EPSCs were made in the absence of GABA_A-R antagonists.

Solubility & Handling

Storage instructions Solubility overview	Room temperature (desiccate) Soluble in water (20mM)
Handling	Hydroscopic solid, contact with air may cause material to change colour and become sticky. Product
	performance should not be affected but we recommend storing the material in a sealed jar.
Important	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not
	for human or veterinary use.

Chemical Data

Chemical name Molecular Weight Chemical structure

Molecular Formula CAS Number PubChem identifier SMILES Source InChi

InChiKey **MDL** number Appearance

NC 2Na¹ O₂N

276.12

6-Cyano-7-nitroquinoxaline-2,3-dione disodium

C₉H₂N₄O₄Na₂ 479347-85-8 2821 C1=C(C(=CC2=C1N=C(C(=N2)[O-])[O-])[N+](=O)[O-])C#N.[Na+].[Na+] Synthetic InChI=1S/C9H4N4O4.2Na/c10-3-4-1-5-6(2-7(4)13(16)17)12-9(15)8(14)11-5;;/h1-2H,(H,11,14)(H,12, 15);;/q;2*+1/p-2 YCXDDPGRZKUGDG-UHFFFAOYSA-L MFCD09953908 Brown or yellow solid

References

6,7-Dinitro-quinoxaline-2,3-dion and 6-nitro,7-cyano-quinoxaline-2,3-dion antagonise responses to NMDA in the rat spinal cord via an action at the strychnine-insensitive glycine receptor.

Birch PJ et al (1988) Eur J Pharmacol 156(1) PubMedID 2905271

The calpain inhibitor MDL-28170 and the AMPA/KA receptor antagonist CNQX inhibit neurofilament degradation and enhance neuronal survival in kainic acid-treated hippocampal slice cultures.

Lopez-Picon FR et al (2006) Eur J Neurosci 23(10) PubMedID 16817871

6-Cyano-7-nitroquinoxaline-2,3-dione (CNQX) increases GABAA receptor-mediated spontaneous postsynaptic currents in the dentate granule cells of rat hippocampal slices.

Hashimoto Y et al (2004) Neurosci Lett 358(1) **PubMedID** 15016428

Pharmacological characterization of glutamatergic agonists and antagonists at recombinant human homomeric and heteromeric kainate receptors in vitro.

Alt et al (2004) Neuropharmacology 46(6) PubMedID 15033339