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%

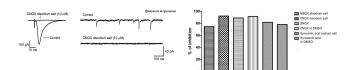
Customer comments The CNQX is going fine ! Verified customer, IBPS, Inserm, CNRS

It works exactly as it should! Dissolved in water, kept in aliquots in -20 freezer. Verified customers,

SickKids (University of Toronto)

Description 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.

CNQX also available.

Application notes

The AMPA receptor antagonist CNQX disodium salt is commonly used at concentrations of 10 μ M 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 μs) 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

Solubility overviewSoluble in water (20mM)Storage instructionsRoom temperature (desiccate)

Storage of solutions Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if

storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.

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.

Shipping Conditions Stable for ambient temperature shipping. Follow storage instructions on receipt.

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 structure

Handling

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

Molecular Weight 276.12

 O_2N

PubChem identifier 2821

SMILES C1=C(C(=CC2=C1N=C(C(=N2)[O-])[O-])[N+](=O)[O-])C#N.[Na+].[Na+]

Source Synthetic

InChi 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

InChiKey YCXDDPGRZKUGDG-UHFFFAOYSA-L

MDL number MFCD09953908
Appearance 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