

## DATASHEET

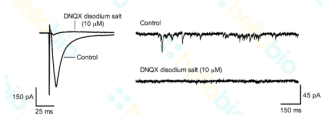
### DNQX disodium salt

## Product overview

<b>Name</b>	DNQX disodium salt
<b>Cat No</b>	HB0262
<b>Description</b>	Selective, competitive AMPA / kainate receptor antagonist. Sodium salt.
<b>Biological action</b>	Antagonist
<b>Purity</b>	>98%
<b>Customer comments</b>	<i>DNQX disodium salt is a good product</i> <b>Verified customer, Research University Paris</b>

## Images

Fig 1: DNQX disodium salt inhibition of evoked and spontaneous glutamate mediated EPSCs in mouse cortical neuron



DNQX disodium salt antagonizes the actions of glutamate at AMPA receptors. It is commonly used to reduce excitatory post synaptic currents (EPSC) and is commonly used at 10 μM. DNQX disodium salt from Hello Bio completely blocks both spontaneous and evoked EPSCs at 10 μM, with concentrations of 1 μM also effective. For assay protocol, see [Protocol 1](#) in Application Notes below.



## Biological Data

### Biological description

DNQX disodium salt is a water soluble, selective and competitive AMPA and kainate receptor antagonist. It also acts as partial AMPA agonist in the presence of  $\gamma 2$  transmembrane AMPA receptor regulatory proteins (TARP) subunit.

DNQX is also a neuroleptic agent that displays pro-oxidant activity.

**DNQX** freebase is also available.

### Application notes

DNQX disodium salt antagonizes the actions of glutamate at AMPA receptors. It is commonly used to reduce excitatory post synaptic currents (EPSC) and is commonly used at 10 μM. DNQX disodium salt from Hello Bio completely blocks both spontaneous and evoked EPSCs at 10 μM, with concentrations of 1 μM also effective (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 prefrontal 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 DNQX disodium salt until complete receptor inhibition.
- Spontaneous EPSCs were recorded before and after addition of DNQX disodium

- salt by holding the neuron at -70 mV and recording for 10 sec.
- Recordings for EPSCs were made in the absence of GABA<sub>A</sub>-R antagonists.

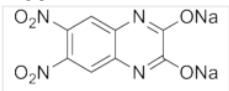
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## Solubility & Handling

<b>Storage instructions</b>	Room temperature (desiccate)
<b>Solubility overview</b>	Soluble in water (100mM)
<b>Important</b>	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

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## Chemical Data

<b>Chemical name</b>	6,7-Dinitroquinoxaline-2,3-dione disodium salt
<b>Molecular Weight</b>	296.1
<b>Chemical structure</b>	
<b>Molecular Formula</b>	C <sub>8</sub> H <sub>2</sub> N <sub>4</sub> Na <sub>2</sub> O <sub>6</sub>
<b>CAS Number</b>	1312992-24-7
<b>PubChem identifier</b>	45073428
<b>SMILES</b>	<chem>C1=C2C(=CC(=C1[N+](=O)[O-])[N+](=O)[O-])N=C(C(=N2)[O-])[O-].[Na+].[Na+]</chem>
<b>Source</b>	Synthetic
<b>InChi</b>	InChI=1S/C8H4N4O6.2Na/c13-7-8(14)10-4-2-6(12(17)18)5(11(15)16)1-3(4)9-7;;/h1-2H,(H,9,13)(H,10,14);;q;2*+1/p-2
<b>InChiKey</b>	GPSBSOYURFUVKJ-UHFFFAOYSA-L
<b>Appearance</b>	Brown solid

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## References

### Redox properties and prooxidant cytotoxicity of a neuroleptic agent 6,7-dinitrodihydroquinoxaline-2,3-dione (DNQX).

Šarlauskas J *et al* (2013) Acta Biochim Pol 60(2)

**PubMedID** [23757451](#)

### TARP auxiliary subunits switch AMPA receptor antagonists into partial agonists.

Menuz K *et al* (2007) Science 318(5851)

**PubMedID** [17975069](#)

### Selective excitatory actions of DNQX and CNQX in rat thalamic neurons.

Lee SH *et al* (2010) J Neurophysiol 103(4)

**PubMedID** [20107128](#)

### 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](#)

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