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DATASHEET

Kainic acid

Product overview

NameKainic acidCat NoHB0355Biological actionAgonistPurity>98%

Customer comments We tested the Kainic acid and we are very happy with it. Hello Bio Kainic acid was used at 25 mg/kg

 $\it i.p.\ and\ successfully\ induced\ status\ epilepticus.\ {\bf Verified\ customer,\ CNRS,\ France}$

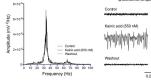
Exactly as ordered - we ordered 50 mg kainic acid and it arrived swiftly. The batch display same EC50 and efficacy as all other batches, so we assume it is pure and an intact batch of the compound. Verified customer, University of Copenhagen

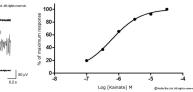
Very happy - service from Hello Bio is always friendly and quick. Kainic acid (KA) works as expected on AMPA receptors in HEK cells. Much cheaper than alternative suppliers. Verified customer, UCL

Description

Prototypic, selective kainate receptor agonist. Potent excitant and neurotoxin.

Images









Biological Data

Biological description

Kainic acid is the prototypic, selective kainate receptor (KAR) agonist. Analog of L-Glutamate.

Kainic acid is a potent excitant and neurotoxin (shows ~30-fold more neurotoxic potency than L-Glutamate).

It induces various changes *in vivo* including recurrent seizures, behavioural changes, oxidative stress, glial activation and selective neuronal death.

Kainic acid exerts its neuroexcitatory action by binding to glutamate receptors to cause Ca²⁺ influx and triggering subsequent excitotoxic neuronal death pathway cascades.

Application notes

It is widely used to model various neurodegenerative diseases and epilepsy.

The kainate receptor (KAR) agonist kainic acid induces neuronal oscillations in the gamma frequency band (35-100Hz). KAR activation leads to synchronous activity of excitatory and inhibitory network firing. Kainic acid is commonly used at concentrations of 400-600 nM to induce oscillations in acute brain slices. At a concentration of 550 nM, kainic acid from Hello Bio induces a strong gamma frequency neuronal oscillation (approx. 35Hz) within the CA3 region of the rat hippocampus (see Fig 1 above). Oscillations were abolished following kainic acid washout.

#Protocol 1: Oscillations assay

- Acute rat hippocampal slices were placed in an interface chamber and held at 30 °C with a glass recording electrode placed in the CA3 region of the hippocampus.
- Control basal activity was recorded for 1 hr in the presence of aCSF before continuous perfusion of aCSF containing kainic acid at 550nM to induce oscillations.
- Oscillation activity was recorded once oscillations had stabilised (~2-3 hours).
- Kainic acid was then washed out and replaced with control aCSF.
- Data were analysed to create a power spectrum of the oscillation.

Solubility & Handling

Solubility overview Soluble in water (25mM, gentle warming)

213.23

Storage instructions Room temperature

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.

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 name (2S,3S,4S)-Carboxy-4-(1-methylethenyl)-3-pyrrolidineacetic acid

Molecular Weight Chemical structure

 CO_2H

SMILES CC(=C)[C@H]1CNC([C@H]1CC(=O)O)C(=O)O

Source Extracted from diginea simplex

InChi InChi=1S/C10H15NO4/c1-5(2)7-4-11-9(10(14)15)6(7)3-8(12)13/h6-7,9,11H,1,3-4H2,2H3,(H,12,13)(

H,14,15)/t6-,7+,9-/m0/s1

InChiKey VLSMHEGGTFMBBZ-OOZYFLPDSA-N

MDL numberMFCD00150833AppearanceWhite solid

References

Kainic acid-mediated excitotoxicity as a model for neurodegeneration.

Wang Q *et al* (2005) Mol Neurobiol 31(1-3) **PubMedID** 15953808

Distinct modulation of the endocannabinoid system upon kainic acid-induced in vivo seizures and in vitro epileptiform bursting.

Fezza F *et al* (2014) Mol Cell Neurosci 62 **PubMedID** 25064144

Identification and characterization of the ligand binding subunit of a kainic acid receptor using monoclonal antibodies and peptide mapping.

Hampson DR *et al* (1989) J Biol Chem 264(22) **PubMedID** 2546953

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