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

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

customercare-usa@hellobio.com



DATASHEET

(RS)-CPP

Product overview

Name(RS)-CPPCat NoHB0036Biological actionAntagonist

Description Potent, selective, competitive NMDA receptor antagonist

Images



Biological Data

Biological description

Potent, selective and competitive NMDA receptor antagonist which reversibly binds to the glutamate binding site. Crosses the blood brain barrier and is active in vivo. Shows various effects (e.g. supresses seizure activity, interferes with addiction paradigms, blocks LTP and LTD and impairs learning and memory). (R)-CPP also available.

Solubility & Handling

Solubility overview Storage instructions Storage of solutions

Shipping Conditions

Important

Soluble in water (100mM) Room temperature (desiccate)

Storage of solutions

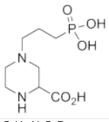
Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.

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

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 (RS)-3-(2-Carboxypiperazin-4-yl)-propyl-1-phosphonic acid 252.21



PubChem identifier 1228

SMILES C1CN(CC(N1)C(=O)O)CCCP(=O)(O)O

InChi InChi=1S/C8H17N2O5P/c11-8(12)7-6-10(4-2-9-7)3-1-5-16(13,14)15/h7,9H,1-6H2,(H,11,12)(H2,13,

14,15)

InChiKey CUVGUPIVTLGRGI-UHFFFAOYSA-N

MDL number MFCD00055136

References

CPP, a new potent and selective NMDA antagonist. Depression of central neuron responses, affinity for [3H]D-AP5 binding sites on brain membranes and anticonvulsant activity.

Davies J *et al* (1986) Brain Res 382(1) **PubMedID**2876749

Action of 3-((+/-)-2-carboxypiperazin-4-yl)-propyl-1-phosphonic acid (CPP): a new and highly potent antagonist of N-methyl-D-aspartate receptors in the hippocampus.

Harris EW *et al* (1986) Brain Res 382(1) **PubMedID**2876750

CPP, a selective N-methyl-D-aspartate (NMDA)-type receptor antagonist: characterization in vitro and in vivo.

Lehmann J et al (1987) J Pharmacol Exp Ther 240(3)

PubMedID 2882014

Measurement of NMDA Receptor Antagonist, CPP, in Mouse Plasma and Brain Tissue Following Systematic Administration Using Ion-Pair LCMS/MS.

Gemperline E et al (2014) Analytical methods: advancing methods and applications 6

PubMedID 25663848