abcam

Product datasheet

MPEP hydrochloride, mGlu5 antagonist ab120008

34 References 4 图像

概述

产品名称 MPEP hydrochloride, mGlu5拮抗剂

描述 Potent, selective mGlu₅拮抗剂

生物学描述 Subtype selective and potent non-competitive mGlu $_5$ antagonist (IC $_{50}$ = 36 nM). Central effects

following systemic administration in vivo.

纯度 > 99%

CAS编号 219911-35-0

化学结构

H₃C N

性能

化学名称 2-Methyl-6-(phenylethynyl)pyridine hydrochloride

分子量 229.71

分子式 C₁₄H₁₁N.HCl

PubChem识别号 9794588

存放说明 Store at +4°C. Store under desiccating conditions. The product can be stored for up to 12

months.

溶解度概述 Soluble in water to 5 mM, in ethanol to 100 mM and in DMSO to 100 mM

处理 Wherever possible, you should prepare and use solutions on the same day. However, if you need

to make up stock solutions in advance, we recommend that you store the solution as aliquots in tightly sealed vials at -20°C. Generally, these will be useable for up to one month. Before use, and

prior to opening the vial we recommend that you allow your product to equilibrate to room

temperature for at least 1 hour.

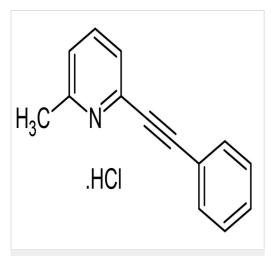
Need more advice on solubility, usage and handling? Please visit our frequently asked

questions (FAQ) page for more details.

SMILES [CI-].Cc2cccc(C#Cc1ccccc1)[nH+]2

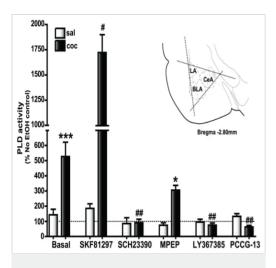
来源 Synthetic

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Chemical Structure - MPEP hydrochloride, $mGlu_5$ antagonist (ab120008)

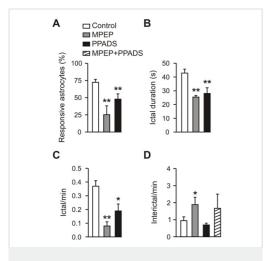
2D chemical structure image of ab120008, MPEP hydrochloride, mGlu5 antagonist



Functional Studies - MPEP hydrochloride, mGlu₅ antagonist (ab120008)

Image from Krishnan B, et al. Plos One, 6(9), e25639. Fig 7,; doi: 10.1371/journal.pone.0025639

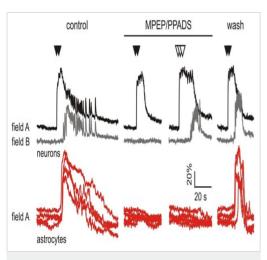
Basal PLD activity is strongly stimulated by the D1/5R agonist and blocked by the D1/5R, mGluR5, mGluR1, and the PLD-linked mGluR antagonists in the amygdala of cocaine CPP animals. The dotted line indicates PLD activity associated with control slices (no EtOH added) which was determined for each animal and used to calculate the change in PLD activity levels with EtOH and/or drug application.



Astrocyte Ca2+ signal inhibition does not affect interictal discharges. (A–D) Mean percentage of astrocytes activated by the ictal discharges (A), mean duration (B) and frequency (C) of the ictal discharge, and mean frequency of interictal discharges (D) under different experimental conditions in EC slice preparations. Controls (n=16), MPEP (ab120008) (n=7), PPADS (ab120009) (n=9), and MPEP+PPADS (n=3). A single asterisk (*) indicates p<0.05; double asterisks (**), p<0.01.

Functional Studies - MPEP hydrochloride, mGlu₅ antagonist (ab120008)

Image from Gómez-Gonzalo Met al., PLoS Biol. 2010;8(4):e1000352. Fig 2.; doi: 10.1371/journal.pbio.1000352. Reproduced under the Creative Commons license http://creativecommons.org/licenses/by/4.0/



Functional Studies - MPEP hydrochloride, m Glu_5

Image from Gómez-Gonzalo Met al., PLoS Biol. 2010;8(4):e1000352. Fig 6(A).; doi: 10.1371/journal.pbio.1000352.

antagonist (ab120008)

Ca2+ signal from a field A neuron, a field B neuron, and field A astrocytes in response to repetitive episodes of NMDA stimulation (black arrowheads). The NMDA stimulation that evoked an ictal discharge became ineffective after blocking the astrocyte response by bath perfusion with MPEP (ab120008) and PPADS (ab120009). An ictal discharge could be recovered by increasing the number of NMDA puffs (white arrowheads). A double NMDA pulse evoked both astrocyte activation and the ictal discharge after inhibitor washout.

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