abcam

Product datasheet

MPEP hydrochloride, mGlu5 antagonist ab120008

34 References 画像数 4

製品の概要

製品名 MPEP hydrochloride, mGlu5 antagonist

製品の詳細 Potent, selective mGlu₅ antagonist

生理活性の詳細 Subtype selective and potent non-competitive mGlu₅ antagonist (IC₅₀ = 36 nM). Central effects

following systemic administration in vivo.

精製度 > 99%

CAS 番号 219911-35-0

構造式

H₃C N .HCl

製品の特性

体系名 2-Methyl-6-(phenylethynyl)pyridine hydrochloride

分子量 229.71

分子式 C₁₄H₁₁N.HCl

PubChem 登録番号 9794588

保存方法 Store at +4°C. Store under desiccating conditions. The product can be stored for up to 12

months.

溶解性 Soluble in water to 5 mM, in ethanol to 100 mM and in DMSO to 100 mM

使用に関する注意 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.

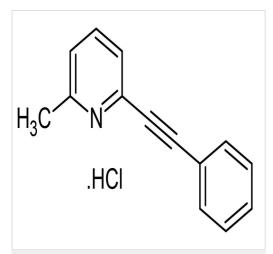
Need more advice on solubility, usage and handling? Please visit our frequently asked

questions (FAQ) page for more details.

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

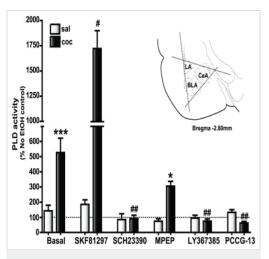
由来 Synthetic

1



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

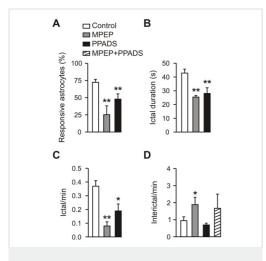
Chemical Structure - MPEP hydrochloride, $mGlu_5$ antagonist (ab120008)



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

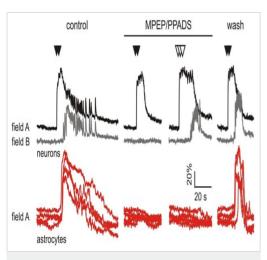
Basal PLD activity is strongly stimulated by the D1/5R agonist and blocked by the D1/5R, mGluR5, mGluR1, and the PLD-linked mGluR antagonists in the amygdala of cocaine CPP animals. The dotted line indicates PLD activity associated with control slices (no EtOH added) which was determined for each animal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