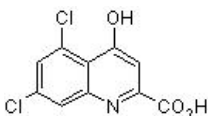


Product datasheet

5,7-Dichlorokynurenic acid, NMDA receptor glycine site antagonist ab120023

2 图像

概述

产品名称	5,7-Dichlorokynurenic acid, NMDA receptor glycine site拮抗剂
描述	NMDA receptor glycine site拮抗剂
生物学描述	Potent NMDA receptor glycine site antagonist. Water soluble form available - see (ab120254).
CAS编号	131123-76-7
化学结构	

性能

化学名称	5,7-Dichloro-4-hydroxyquinoline-2-carboxylic acid
分子量	258.06
分子式	C ₁₀ H ₅ Cl ₂ NO ₃
PubChem识别号	1779
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 1 eq. NaOH to 50 mM
处理	<p>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.</p> <p>Refer to SDS for further information</p> <p>Need more advice on solubility, usage and handling? Please visit our frequently asked questions (FAQ) page for more details.</p>
SMILES	<chem>O=C(O)c1cc(O)c2c(Cl)cc(Cl)cc2n1</chem>
来源	Synthetic

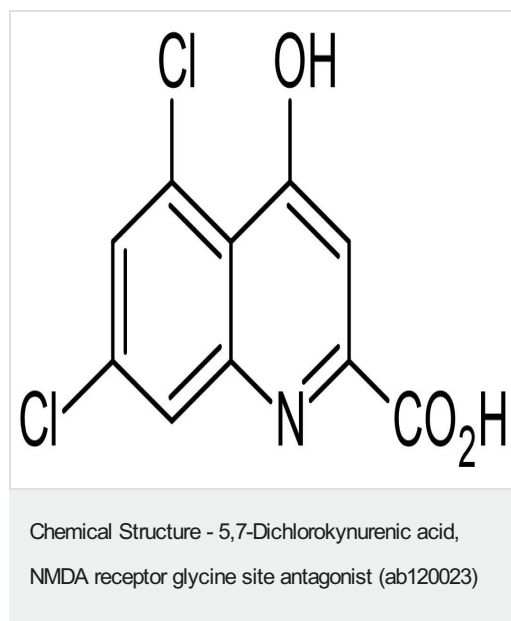
The Abpromise guarantee

Abpromise™承诺保证使用ab120023于以下的经测试应用

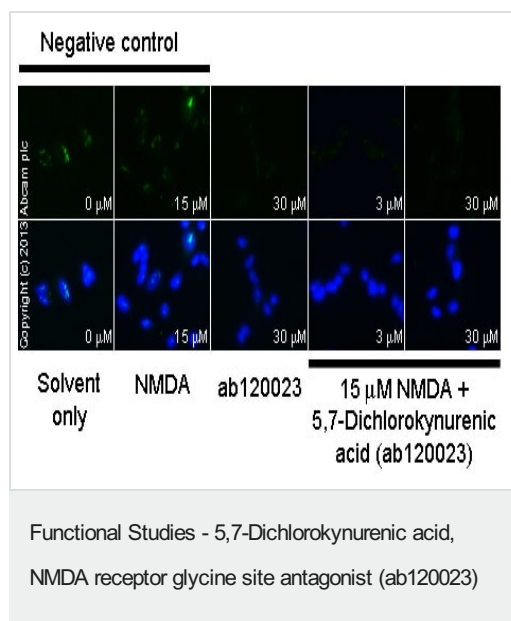
“应用说明”部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Functional Studies		Use at an assay dependent concentration.

图片



2D chemical structure image of ab120023, 5,7-Dichlorokynurenic acid, NMDA receptor glycine site antagonist



ab12416 staining cGMP in SKNSH cells treated with 5,7-Dichlorokynurenic acid (ab120023), by ICC/IF. Decrease in cGMP expression correlates with increased concentration of 5,7-Dichlorokynurenic acid, as described in literature.

The cells were incubated at 37°C for 20 minutes in media containing different concentrations of ab120023 (5,7-Dichlorokynurenic acid) in DMSO. Some samples were then further incubated with 15 μM NMDA (**ab120052**) for 5 minutes and all samples were fixed with 100% methanol for 5 minutes at -20°C and blocked with PBS containing 10% goat serum, 0.3 M glycine, 1% BSA and 0.1% tween for 2h at room temperature. Staining of the treated cells with **ab12416** (5 μg/ml) was performed overnight at 4°C in PBS containing 1% BSA and 0.1% tween. A DyLight 488 anti-rabbit polyclonal antibody (**ab96899**) at 1/250 dilution was used as the secondary antibody. Nuclei were counterstained with DAPI and are shown in blue.

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