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Certificate of Analysis

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Print Date: Oct 11th 2024

Product Name: (S)-3,4-DCPG

201730-11-2 CAS Number: IUPAC Name: (S)-3,4-Dicarboxyphenylglycine

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: Batch Molecular Weight: Physical Appearance: Solubility:

Storage: **Batch Molecular Structure:**

C10H9NO6.1/2H2O 248.19 White solid water to 100 mM DMSO to 25 mM Desiccate at RT



2. ANALYTICAL DATA

Chiral HPLC: Shows 99.2% purity ¹H NMR: Consistent with structure Mass Spectrum: Consistent with structure $[\alpha]_D$ = +122.3 (Concentration = 0.5, Solvent = 6M HCl) **Optical Rotation: Microanalysis:** Carbon Hydrogen Nitrogen Theoretical 48.39 4.06 5.64 47.63 Found 4.07 5.5

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

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Catalog No.: 1302 Batch No.: 12

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Product Name: (S)-3,4-DCPG

CAS Number: 201730-11-2

IUPAC Name: (S)-3,4-Dicarboxyphenylglycine

Description:

(S)-3,4-DCPG is a potent, selective mGlu_{8a} agonist (EC₅₀ = 31 nM). Displays > 100-fold selectivity over mGlu₁₋₇ and displays little or no activity at NMDA and kainate receptors. Increases c-Fos expression in stress-related brain areas following systemic administration in mice in vivo. Also potent anticonvulsant in mice in vivo. Caged (S)-3,4-DCPG, Racemate and R-enantiomer also available.

Physical and Chemical Properties:

Batch Molecular Formula: C₁₀H₉NO₆.½H₂O Batch Molecular Weight: 248.19 Physical Appearance: White solid

Minimum Purity: ≥98%

Batch Molecular Structure:



References:

Linden *et al* (2003) Systemic administration of the potent mGlu8 receptor agonist (S)-3,4-DCPG induces c-Fos in stress-related brain regions in wild-type, but not mGlu8 receptor knockout mice. Neuropharmacology **45** 473. PMID: 12907308.

Moldrich *et al* (2001) Anticonvulsant activity of 3,4-dicarboxyphenylglycines in DBA/2 mice. Neuropharmacology **40** 732. PMID: 11311902.

Thomas *et al* (2001) (S)-3,4-DCPG, a potent and selective mGlu8a receptor agonist activates metabotropic glutamate receptors on primary afferent terminals in the neonatal rat spinal cord. Neuropharmacology **40** 311. PMID: 11166323.

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Catalog No.: 1302

12

Storage: Desiccate at RT

Solubility & Usage Info:

water to 100 mM DMSO to 25 mM

CAUTION - Analysis shows that this material rapidly decomposes when dissolved in alkaline solution. Therefore we recommend that this product is dissolved in water.

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. *Unless contradicted by product-specific protocols or instructions, our standard recommendations apply:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.