

Product Name: L-*trans*-2,4-PDC

Catalog No.: 0298

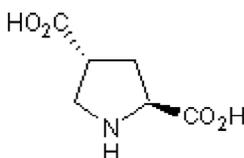
Batch No.: 15

CAS Number: 64769-66-0

IUPAC Name: L-*trans*-Pyrrolidine-2,4-dicarboxylic acid

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: C₆H₉NO₄
Batch Molecular Weight: 159.14
Physical Appearance: White solid
Solubility: water to 100 mM
 phosphate buffered saline to 100 mM
 1eq. NaOH to 100 mM
Storage: Store at RT
Batch Molecular Structure:



2. ANALYTICAL DATA

HPLC: Shows >99.4% purity
¹H NMR: Consistent with structure
Mass Spectrum: Consistent with structure
Optical Rotation: [α]_D = -52 (Concentration = 1, Solvent = Water)
Microanalysis:

	Carbon	Hydrogen	Nitrogen
Theoretical	45.28	5.7	8.8
Found	44.98	5.73	8.64

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

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CAS Number: 64769-66-0

IUPAC Name: L-*trans*-Pyrrolidine-2,4-dicarboxylic acid

Description:

L-*trans*-2,4-PDC is a potent, competitive, transportable EAAT1-4 inhibitor/non-transportable EAAT5 inhibitor. In [³H]-d-Asp uptake assays in HEK293 cells expressing human EAAT1, EAAT2 and EAAT3, K_i values are 20, 20 and 109 μM, respectively. In a FLIPR Membrane Potential (FMP) assay, K_m values for L-*trans*-2,4-PDC are 7.7, 11 and 19 μM for human EAAT2, EAAT3 and EAAT1, respectively.

Physical and Chemical Properties:

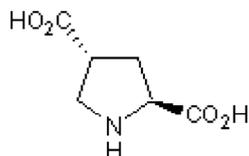
Batch Molecular Formula: C₆H₉NO₄

Batch Molecular Weight: 159.14

Physical Appearance: White solid

Minimum Purity: ≥97%

Batch Molecular Structure:



References:

Jensen and Bräuner-Osborne (2004) Pharmacological characterization of human excitatory amino acid transporters EAAT1, EAAT2 and EAAT3 in a fluorescence-based membrane potential assay. *Biochem.Pharmacol.* **67** 2115. PMID: 15135308.

Zuiderwijk et al (1996) Effects of uptake carrier blockers SK & F 89976-A and L-*trans*-PDC on *in vivo* release of amino acids in rat hippocampus. *Eur.J.Pharmacol.* **307** 275. PMID: 8836615.

Mitrovic and Johnston (1994) Regional differences in the inhibition of L-glutamate and L-aspartate sodium-dependent high affinity uptake systems in rat CNS synaptosomes by L-*trans*-pyrrolidine-2,4-dicarboxylic *threo*-3-hydroxy-D-aspartate and D-aspartate. *Neurochem.Int.* **24** 583. PMID: 7981641.

Storage: Store at RT

Solubility & Usage Info:

water to 100 mM

phosphate buffered saline to 100 mM

1eq. NaOH to 100 mM

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.

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