

Certificate of Analysis

Print Date: Jan 15th 2016

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Product Name: NPS 2390 Catalog No.: 4134 Batch No.: 1

CAS Number: 226878-01-9

IUPAC Name: N-tricyclo[3.3.1.13,7]dec-1-yl-2-quinoxalinecarboxamide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: $C_{19}H_{21}N_3O$ Batch Molecular Weight:307.39Physical Appearance:Grey solid

Solubility: DMSO to 50 mM

ethanol to 50 mM

Storage: Store at RT

Batch Molecular Structure:

2. ANALYTICAL DATA

TLC: $R_f = 0.37$ (Chloroform:Methanol [95:5])

HPLC: Shows 99.9% purity

¹H NMR: Consistent with structure

Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 74.24 6.89 13.67 Found 74.15 6.95 13.75



Product Information

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Description:

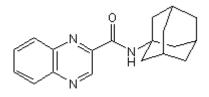
Group I mGlu antagonist; displays noncompetitive antagonist activity at both mGlu1 and mGlu5 receptors. Thought to act on a site separate from the glutamate binding pocket.

Physical and Chemical Properties:

Batch Molecular Formula: C₁₉H₂₁N₃O Batch Molecular Weight: 307.39 Physical Appearance: Grey solid

Minimum Purity: >99%

Batch Molecular Structure:



Storage: Store at RT

Solubility & Usage Info:

DMSO to 50 mM ethanol to 50 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. Our standard recommendations are:

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.

References:

van Wagenen et al (2000) Structure-activity relationship studies of NPS 2390: a potent and selective group I metabotropic glutamate receptor antagonist. Abstr. Soc. Neurosci. Program No. 618.3..

Layreysen et al (2003) [3H]R214127: a novel high-affinity radioligand for the mGlu1 receptor reveals a common binding site shared by multiple allosteric antagonists. Mol. Pharmacol. 63 1082. PMID: 12695537.

Mabire et al (2005) Synthesis, structure-activity relationship, and receptor pharmacology of a new series of quinoline derivatives acting as selective, noncompetitive mGlu1 antagonists. J.Med.Chem. 48 2134. PMID: 15771457.