

Certificate of Analysis

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Print Date: Jan 15th 2016

Product Name: PF 750 Catalog No.: 3307 Batch No.: 1

CAS Number: 959151-50-9

IUPAC Name: N-Phenyl-4-(3-quinolinylmethyl)-1-piperidinecarboxamide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: $C_{22}H_{23}N_3O.34H_2O$

Batch Molecular Weight: 358.95

Physical Appearance: Off-white solid

Solubility: DMSO to 100 mM ethanol to 100 mM

Storage: Store at RT

Batch Molecular Structure:

2. ANALYTICAL DATA

TLC: $R_f = 0.62$ (Ethyl acetate:Methanol [95:5])

HPLC: Shows 98% purity

¹H NMR: Consistent with structure Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 73.61 6.88 11.71 Found 73.57 6.63 11.79



Product Information

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Description:

Irreversible fatty acid amide hydrolase (FAAH) inhibitor (IC $_{50}$ = 16.2 nM) that displays no activity at a range of other serine hydrolases. Selectively inhibits FAAH within the central nervous system. Orally active.

Physical and Chemical Properties:

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Batch Molecular Weight: 358.95 Physical Appearance: Off-white solid

Minimum Purity: >98%

Batch Molecular Structure:

Storage: Store at RT

Solubility & Usage Info:

DMSO to 100 mM ethanol 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. 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:

Ahn et al (2007) Novel mechanistic class of fatty acid amide hydrolase inhibitors with remarkable selectivity. Biochemistry 46 13019. PMID: 17949010.

Mileni et al (2008) Structure-guided inhibitor design for human FAAH by interspecies active site conversion. Proc.Natl.Acad.Sci.USA 105 12820.