TOCRIS b i o s c i e n c e

Certificate of Analysis

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Print Date: Jan 30th 2014

Product Name: SRPIN340

Catalog No.: 5063 Batch No.: 1

CAS Number: 218156-96-8

IUPAC Name: N-[2-(1-Piperidinyl)-5-(trifluoromethyl)phenyl]-4-pyridinecarboxamide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: Batch Molecular Weight: Physical Appearance: Solubility: C₁₈H₁₈F₃N₃O 349.35 Off-white solid DMSO to 100 mM 1eq. HCl to 50 mM ethanol to 100 mM Store at +4°C

Storage: Batch Molecular Structure:

OF3 N N N

2. ANALYTICAL DATA

HPLC: ¹H NMR: Mass Spectrum: Microanalysis: Shows 99.8% purity Consistent with structure Consistent with structure

	Carbon	Hydrogen	Nitrogen
Theoretical	61.88	5.19	12.03
Found	61.92	5.08	11.99

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

 Corris Bioscience is an R&D Systems company

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218156-96-8

Product Information

Storage: Store at +4°C

DMSO to 100 mM

1eg. HCl to 50 mM

ethanol to 100 mM

a 45-60°C water bath).

are:

Solubility & Usage Info:

Stability and Solubility Advice:

for up to 6 months from date of receipt.

Some solutions can be difficult to obtain and can be

encouraged by rapid stirring, sonication or gentle warming (in

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

SOLIDS: Provided storage is as stated on the product label

and the vial is kept tightly sealed, the product can be stored

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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CAS Number:

IUPAC Name: N-[2-(1-Piperidinyl)-5-(trifluoromethyl)phenyl]-4-pyridinecarboxamide

Description:

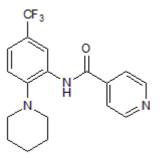
Selective serine arginine protein kinase (SRPK) inhibitor for SRPK1 (K_i value = 0.89 μ M). Also inhibits SRPK2. Does not significantly inhibit other SRPKs such as CLK1 and CLK4, or other classes of SR kinases. Suppresses choroidal neovascularization formation growth. Reduces the expression of proangiogenic VEGF165 without affecting VEGF165b expression. Suppresses RNA virus Sindbis propagation (IC₅₀ = 60 μ M in plaque assays in Vero cultures) and HCV-JFH1 replication in Huh7.5.1 cultures.

Physical and Chemical Properties:

Batch Molecular Formula: C₁₈H₁₈F₃N₃O Batch Molecular Weight: 349.35 Physical Appearance: Off-white solid

Minimum Purity: >98%

Batch Molecular Structure:



References:

Fukuhara *et al* (2006) Utilization of host SR protein kinases and RNA-splicing machinery during viral replication. Proc.Natl.Acad.Sci.USA **103** (30) 11329. PMID: 16840555.

Karakama *et al* (2010) Inhibition of hepatitis C virus replication by a specific inhibitor of serine-arginine-rich protein kinase. Antimicrob.Agents.Chemother. **54** (8) 3179. PMID: 20498328.

Gammons *et al* (2013) SRPK1 inhibition modulates VEGF splicing to reduce pathological neovascularization in a rat model of retinopathy of prematurity. Invest.Ophthalmol.Vis.Sci. **54** (8) 5797. PMID: 23761094.

Dong *et al* (2013) Specific inhibition of serine/arginine-rich protein kinase attenuates choroidal neovascularization. Mol.Vis. **19** 536. PMID: 23559848.

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