

# **Certificate of Analysis**

Print Date: Dec 7th 2022

www.tocris.com

Product Name: FK 866 hydrochloride Catalog No.: 4808 Batch No.: 2

CAS Number: 2727965-45-7

IUPAC Name: 2-(E)-N-[4-(1-Benzoyl-4-piperidinyl)butyl]-3-(3-pyridinyl)-2-propenamide hydrochloride

#### 1. PHYSICAL AND CHEMICAL PROPERTIES

**Batch Molecular Formula:**  $C_{24}H_{29}N_3O_2.HCI.\frac{1}{4}H_2O$ 

Batch Molecular Weight: 432.47

Physical Appearance: Off-white solid

**Solubility:** water to 5 mM with gentle warming

DMSO to 100 mM

Storage: Store at -20°C

**Batch Molecular Structure:** 

#### 2. ANALYTICAL DATA

**TLC:**  $R_f = 0.44$  (Dichloromethane:Methanol [9:1])

HPLC: Shows >99.9% purity

<sup>1</sup>H NMR: Consistent with structure

Mass Spectrum: Consistent with structure

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 66.65 7.11 9.72 Found 66.75 7.32 9.73

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



# **Product Information**

Print Date: Dec 7th 2022

2

www.tocris.com

Product Name: FK 866 hydrochloride

CAS Number: 2727965-45-7

IUPAC Name: 2-(E)-N-[4-(1-Benzoyl-4-piperidinyl)butyl]-3-(3-pyridinyl)-2-propenamide hydrochloride

#### **Description:**

FK 866 hydrochloride is a non-competitive and potent inhibitor of NAMPT (nicotinamide phosphoribosyltransferase, PBEF1) ( $K_i$  = 0.3 nM); inhibits NAD biosynthesis. Induces delayed cell death by apoptosis in HepG2 human liver carcinoma cells ( $IC_{50} \sim 1$  nM). Induces apoptosis in four different neuroblastoma cell lines; also induces autophagy in SH-SY5Y cells. Potentiates the cytotoxic effects induced by etoposide (Cat. No. 1226) and cisplatin (Cat. No. 2251).

# **Physical and Chemical Properties:**

Batch Molecular Formula: C24H29N3O2.HCI.1/4H2O

Batch Molecular Weight: 432.47 Physical Appearance: Off-white solid

Minimum Purity: ≥98%

#### **Batch Molecular Structure:**

**Storage:** Store at -20°C. This product is packaged under an inert atmosphere.

Catalog No.: 4808

# Solubility & Usage Info:

water to 5 mM with gentle warming DMSO to 100 mM

Standard retail vials are prepared by lyophilisation. The product may appear as a solid, a gel or a film. Solutions should be made by adding solvent directly to the vial. The vial should then be vortexed vigorously to ensure the product has completely dissolved

#### 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:

**Travelli** *et al* (2011) Reciprocal potentiation of the antitumoral activities of FK866, an inhibitor of nicotinamide phosphoribosyltransferase, and etop. or cisp. in neuroblastoma cells. J.Pharmacol.Exp.Ther. *338* 829. PMID: 21685314.

**Galli** *et al* (2008) Synthesis and biological evaluation of isosteric analogues of FK866, an inhibitor of NAD salvage. ChemMedChem **3** 771. PMID: 18247435.

**Hasmann** *et al* (2003) FK866, a highly specific noncompetitive inhibitor of nicotinamide phosphoribosyltransferase, represents a novel mechanism for induction of tumor cell apoptosis. Cancer Res. *63* 7436. PMID: 14612543.

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