



# **Certificate of Analysis**

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Product Name: Indirubin-3'-oxime Catalog No.: 1813 Batch No.: 1

CAS Number: 160807-49-8

IUPAC Name: 3-[1,3-Dihydro-3-(hydroxyimino)-2*H*-indol-2-ylidene]-1,3-dihydro-2*H*-indol-2-one

### 1. PHYSICAL AND CHEMICAL PROPERTIES

**Batch Molecular Formula:**  $C_{16}H_{11}N_3O_2.^3/4H_2O$ 

Batch Molecular Weight: 290.79
Physical Appearance: Red solid

Solubility: DMSO to 100 mM

ethanol to 100 mM

Storage: Store at RT

**Batch Molecular Structure:** 

### 2. ANALYTICAL DATA

**TLC:**  $R_f = 0.58$  (Dichloromethane:Methanol [10:1])

Melting Point:Greater than 250°CHPLC:Shows 99.3% purity

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

Microanalysis: Carbon Hydrogen Nitrogen

Theoretical 66.09 4.33 14.45 Found 66.29 4.2 14.78

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



# **Product Information**

Print Date: Jul 4th 2019

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#### **Description:**

Inhibitor of GSK-3 $\beta$  (IC<sub>50</sub> = 190 nM) and cyclin dependent kinases (cdks; IC<sub>50</sub> values are 0.05, 0.18 and 0.25-0.5  $\beta$ M for cdk9, cdk1 and cdk2, respectively). Also reported to inhibit AMPK, LCK and SGK. Inhibits cdk5- and GSK-3 $\beta$ -mediated tau phosphorylation. Induces cell cycle arrest and inhibits cell proliferation.

#### **Physical and Chemical Properties:**

Batch Molecular Formula: C<sub>16</sub>H<sub>11</sub>N<sub>3</sub>O<sub>2</sub>.<sup>3</sup>/<sub>4</sub>H<sub>2</sub>O

Batch Molecular Weight: 290.79 Physical Appearance: Red solid

**Minimum Purity:** >99%

## **Batch Molecular Structure:**

#### Storage: Store at RT

CAUTION - This product is light sensitive and we recommend that the solid material and any solutions obtained are protected from exposure to light.

# 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:

Jorda et al (2018) How selective are pharmacological inhibitors of cell-cycle-regulating cyclin-dependent kinases? J.Med.Chem. 61 9105. PMID: 30234987.

Bain et al (2003) The specificities of protein kinase inhibitors: an update. Biochem.J. 371 199. PMID: 12534346.

**Leclerc** *et al* (2001) Indirubins inhibit glycogen synthase kinase-3β and CDK5/p25, two protein kinases involved in abnormal tau phosphorylation in Alzheimer's disease. J.Biol.Chem. **276** 251. PMID: 11013232.

**Hoessel** *et al* (1999) Indirubin, the active constituent of a Chinese antileukaemia medicine, inhibits cyclin-dependent kinases. Nat.Cell.Biol. *1* 60. PMID: 10559866.