



Nevirapine

Catalog No: tcsc2252

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Available Sizes

Size: 10mg

Size: 50mg

Size: 100mg

Size: 200mg



Specifications

CAS No:

129618-40-2

Formula:

 $C_{15}H_{14}N_4O$

Pathway:

Anti-infection; Anti-infection

Target:

Reverse Transcriptase; HIV

Purity / Grade:

>98%

Solubility:

DMSO: 14.29 mg/mL (53.66 mM; Need ultrasonic)

Alternative Names:

BI-RG 587;NSC 641530;NVP

Observed Molecular Weight:

266.3





Product Description

Nevirapine is a non-nucleoside inhibitor of **HIV-1** reverse transcriptase used to treat and prevent HIV/AIDS; with a $\mathbf{K_i}$ of 270 μ M.

IC50 & Target: Ki: 270 μM (HIV-1 reverse transcriptase)^[1]

In Vitro: Nevirapine itself is an inhibitor of only CYP3A4 at concentrations that are well above those of therapeutic relevance (K_i =270 μ M)^[1]. Nevirapine has been used as a re-differentiation agent to treat cancers in several human cancer models. At all doses (100, 200, 350, 500 μ M) tested, nevirapine significantly inhibits cell proliferation after 48 h treatment. At high dose (500 μ M), nevirapine significantly increases the percentage of apoptotic cells compared with control^[2]. Nevirapine is a potent and selective inhibitor (IC_{50} =10-100 nM) of the replication of a wide variety of HIV-1 strains in several cellular assays^[3].

In Vivo: Nevirapine is available for use in combination with nucleoside HIV-1 reverse transcriptase inhibitors (e.g., zidovudine, didanosine, etc.). Nevirapine has received FDA approval for use in combination with HIV-1 protease inhibitors (e.g., saquinavir, ritonavir, indinavir, etc.). In humans, nevirapine is eliminated primarily in the urine as glucuronide conjugates of 2-, 3-, 8-, and 12-hydroxynevirapine^[1]. Nevirapine is completely absorbed in both sexes of mouse, rat, rabbit, monkey, and chimpanzee. Nevirapine is extensively metabolized in both sexes of all animal species studied^[4]. Nevirapine (9 mg/kg, 18 mg/kg and 36 mg/kg) shows significant reduction in ulcer severity score and ulcer index as compared to the control^[5]

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