Web: www.taiclone.com
Tel: +886-2-2735-9682
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# Nelfinavir (Mesylate)

## Catalog No: tcsc0923

### Available Sizes

**Size:** 5mg

**Size:** 10mg

**Size:** 50mg

**Size:** 100mg

**Size:** 200mg

**Size:** 500mg

**Size:** 1g

**Size:** 2g

### Specifications

**CAS No:**
159989-65-8

**Formula:**
$C_{33}H_{49}N_3O_7S_2$

**Pathway:**
Metabolic Enzyme/Protease;Anti-infection

**Target:**
HIV Protease;HIV

**Purity / Grade:**
>98%

**Solubility:**

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DMSO : ≥ 25 mg/mL (37.66 mM)

**Alternative Names:**
AG 1343 Mesylate

**Observed Molecular Weight:**
663.89

## Product Description

Nelfinavir(AG-1341) is a potent and orally bioavailable human immunodeficiency virus HIV-1 protease inhibitor (Ki=2 nM) and is widely prescribed in combination with HIV reverse transcriptase inhibitors for the treatment of HIV infection.

IC50 & Target: Ki: 2 nM (HIV-1 protease)[2].

***In Vitro:*** In vitro exposure (72 hours) of HAECs to NEL (0.25-2 μg/mL) decreased both basal (2.5-fold) and insulin-induced NO production (4- to 5-fold). NEL suppressed insulin-induced phosphorylation of both Akt and eNOS at serine residues 473 and 1177, respectively. NEL decreased tyrosine phosphorylation of IR-β, IRS-1, and PI3K. Coexposure to troglitazone (TRO; 250 nM) ameliorated the suppressive effects of NEL on insulin signaling and NO production. Coexposure to TRO also increased eNOS expression in NEL-treated HAECs[1]. Nelfinavir(AG-1341) is a potent enzyme inhibitor ($K_i$=2 nM) and antiviral agent (HIV-1 $ED_{50}$=14 nM). An X-ray cocrystal structure of the enzyme-Nelfinavir(AG-1341) complex reveals how the novel thiophenyl ether and phenol-amide substituents of the inhibitor interact with the S1 and S2 subsites of HIV-1 protease, respectively[2].

***In Vivo:*** In vivo studies indicate that Nelfinavir(AG-1341) is well absorbed orally in a variety of species and possesses favorable pharmacokinetic properties in humans[2].