Web: www.taiclone.com
Tel: +886-2-2735-9682
Email: order@taiclone.com

NVP-CGM097 (sulfate)

Catalog No: tcsc6873

Available Sizes

Size: 2mg

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg

Specifications

CAS No:
1313367-56-4

Formula:
$C_{38}H_{49}ClN_4O_8S$

Pathway:
Apoptosis

Target:
MDM-2/p53

Purity / Grade:
>98%

Solubility:
H2O : ≥ 140 mg/mL (184.86 mM)

Alternative Names:
CGM097 sulfate

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Observed Molecular Weight:
757.34

Product Description

NVP-CGM097 sulfate is a potent and selective **MDM2** inhibitor with **IC_{50}** of 1.7±0.1 nM for **hMDM2**.

IC50 & Target: IC50 & Target: IC50: 1.7±0.1 nM (hMDM2)[1]

In Vitro: NVP-CGM097 binds to human MDM2 with an IC_{50} of 1.7 nM and shows high selectivity over MDM4 (IC_{50}=2000 nM). NVP-CGM097 is about four times more potent than Nutlin-3a (IC_{50}=8 nM). In addition, NVP-CGM097 shows no significant activity against Bcl-2:Bak, Bcl-2:Bad, Mcl-1:Bak, Mcl-1:NOXA, XIAP:BIR3, and c-IAP:BIR3 protein-protein interactions. NVP-CGM097 significantly inhibits the proliferation of cells expressing wild-type p53, while sparing the p53 null cells with a 35-58-fold difference. NVP-CGM097 is able to significantly redistribute wild-type p53 into the cell nucleus with an IC_{50} of 0.224 μM, demonstrating its ability to inhibit the p53:MDM2 interaction in living cells. NVP-CGM097 significantly inhibits the proliferation of cells expressing wild-type p53, while sparing the p53 null cells with a 35-58-fold difference. NVP-CGM097 inhibtis HCT116 ($p53^{WT/WT}$) with IC_{50} of 454±136 nM[1].

In Vivo: NVP-CGM097 is able to inhibit the interaction between p53 and MDM2 and reactivate the p53 pathway in a MDM2-amplified SJSA-1 human tumor model, as judged by elevation of p21 mRNA levels, a pharmacodynamic (PD) indicator for p53 activity. p21 mRNA levels are found to increase concomitantly with levels of NVP-CGM097 in tumor-bearing rats dosed at 30 mg/kg. The PD response is biphasic and prolonged up to 24 h. Additional p53 target genes such as MDM2 and PUMA mRNA levels are assessed in the tumor samples as well and showed a similar behavior. Daily treatment with NVP-CGM097 dose dependently and significantly inhibits SJSA-1 tumor growth in rats. It promotes stable disease at 20 mg/kg, which is associated with a plasma AUC_{0-24} of 163 μM•h. After iv administration, the total blood clearance (CL) of NVP-CGM097 is 5 mL/min per kg for mouse, 7 mL/min per kg for rat, 3 mL/min per kg for dog, and 4 mL/min per kg for monkey. The apparent terminal half-life ($t_{1/2}$) is long in rodents and monkey (6-12 h) but is comparatively longer in dogs (20 h). After oral dosing, NVP-CGM097 is well absorbed with T_{max} occurring between 1 and 4.5 h in all species tested[1].