

Ro 31-8220 (mesylate)

Catalog No: tcsc1677

Available Sizes

Size: 10mg

Size: 50mg

Specifications

CAS No: 138489-18-6

Formula:

 $C_{26}H_{27}N_5O_5S_2$

Pathway: TGF-beta/Smad;Epigenetics

Target:

PKC;PKC

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 50 mg/mL (90.31 mM); H2O :

Alternative Names:

Ro 31-8220 methanesulfonate; BisindolyImaleimide IX mesylate

Observed Molecular Weight:

553.65

Product Description

Ro 31-8220 mesylate is a potent **PKC** inhibitor, with **IC**₅₀s of 5, 24, 14, 27, 24 and 23 nM for PKCα, PKCβI, PKCβI, PKCγ, PKCε and rat brain PKC, respectively. Ro 31-8220 also significantly inhibits MAPKAP-K1b, MSK1, S6K1 and GSK3β (**IC**₅₀s, 3, 8, 15, and 38 nM,



respectively), with no effect on MKK3, MKK4, MKK6 and MKK7.

IC50 & Target: IC50: 5 nM (PKCα), 24 nM (PKCβI), 14 nM (PKCβII), 27 nM (PKCγ), 24 nM (PKCε), 23 nM (Rat brain PKC)^[1], 3 nM (MAPKAP-K1b), 8 nM (MSK1), 15 nM (S6K1), 38 nM (GSK3β)^[2]

In Vitro: Ro 31-8220 mesylate is a potent PKC inhibitor, with IC₅₀s of 5, 24, 14, 27, 24 and 23 nM for PKC α , PKC β I, PKC β II, PKC γ , PKC ϵ and rat brain PKC, respectively^[1]. Ro 31-8220 also significantly inhibits MAPKAP-K1b, MSK1, S6K1 and GSK3 β (IC₅₀s, 3, 8, 15, and 38 nM, respectively), with no effect on MKK3, MKK4, MKK6 and MKK7. Moreover, Ro 31-8220 directly suppresses voltage-dependent Na⁺ channels^[2]. Ro 31-8220 (1 μ M) is neuroprotective against paraoxon-induced neuronal cell death in cerebellar granule neurons, blocks paraoxon-induced caspase-3 activity, and reduces the paraoxon-induced increase in phospho-PKC pan levels [3].

In Vivo: Ro 31-8220 (6 mg/kg/d, s.c.) is well tolerated, and has half-life of 5.7 hours in mice. Ro 31-8220-treated $MLP^{-/-}$ mice show a dramatic rescue in fractional shortening after treatment for 6 weeks, but the WT mice shows no change^[4].



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