

MK591 (free acid)

Catalog No: tcsc0953

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Size: 100mg

Specifications

CAS No:

136668-42-3

Formula:

 $\mathsf{C}_{\mathsf{34}}\mathsf{H}_{\mathsf{35}}\mathsf{CIN}_{\mathsf{2}}\mathsf{O}_{\mathsf{3}}\mathsf{S}$

Pathway: Immunology/Inflammation

Target:

FLAP

Purity / Grade:

>98%

Solubility:

H2O :

Alternative Names:

Quiflapon

Observed Molecular Weight:

587.17

Copyright 2021 Taiclone Biotech Corp.



Product Description

MK591 (free acid) is a selective and specific 5-Lipoxygenase-activating protein (FLAP) inhibitor with an IC₅₀ value of 1.6 nM in a FLAP binding assay.

IC50 & Target: IC50 value: 1.6 nM (FLAP)^[1].

In Vitro: MK591 (free acid) is a potent inhibitor of leukotriene (LT) biosynthesis in intact human and elicited rat polymorphonuclear leukocytes (PMNLs) (IC₅₀ values 3.1 and 6.1 nM, respectively) and in human, squirrel monkey, and rat whole blood (IC₅₀ values 510, 69, and 9 nM, respectively). MK591 (free acid) has no effect on rat 5-lipoxygenase. MK591 (free acid) has a high affinity for 5-lipoxygenase activating protein (FLAP) as evidenced by an IC₅₀ value of 1.6 nM in a FLAP binding assay and inhibition of the photoaffinity labelling of FLAP by two different photoaffinity ligands. Inhibition of activation of 5-lipoxygenase was shown through inhibition of the translocation of the enzyme from the cytosol to the membrane in human PMNLs^[1].

In Vivo: MK591 (free acid) is a potent inhibitor of LT biosynthesis in vivo, first, following ex vivo challenge of blood obtained from treated rats and squirrel monkeys, second, in a rat pleurisy model, and, third, as monitored by inhibition of the urinary excretion of LTE4 in antigen-challenged allergic sheep. Inhibition of antigen-induced bronchoconstriction by MK591 (free acid) is observed in inbred rats pretreated with methysergide, Ascaris-challenged squirrel monkeys, and Ascaris-challenged sheep (early and late phase response) [1]. Pups were treated with either vehicle or MK591 (free acid) 10, 20, or 40 mg/kg subcutaneously daily for days 1-4, 5-9, or 10-14. On day 14, the lungs were inflated, fixed, and stained for histopathological and morphometric analyses. Hyperoxia groups treated with MK-0591 (free acid) untreated hyperoxia groups showed definite evidence of aberrant alveolarization but no inflammation^[2].



All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!

Copyright 2021 Taiclone Biotech Corp.