

## FK 3311

Catalog No: tcsc1360

Available Sizes

Size: 10mg

Size: 50mg

**Specifications** 

**CAS No:** 116686-15-8

Formula:

 $\mathsf{C}_{15}\mathsf{H}_{13}\mathsf{F}_2\mathsf{NO}_4\mathsf{S}$ 

Pathway: Immunology/Inflammation

## **Target:**

COX

Purity / Grade:

>98%

**Solubility:** 10 mM in DMSO

## **Alternative Names:**

COX-2 Inhibitor V

**Observed Molecular Weight:** 

341.33

## **Product Description**

FK 3311 is a selective inhibitor of COX-2; antiinflammatory agent.

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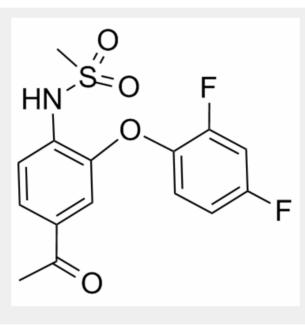
IC50 Value: 1.6 uM [1]

Target: COX-2

Cyclooxygenase (COX) is an intracellular enzyme that converts arachidonic acid into prostaglandin (PG)G2 and PGH2.

in vitro: The racemic mixtures and the (R)- and (S)-isomers of the 2 metabolites were inactive in the PGE2 test. IC50 values were more than 100 uM for (2 and 5), compared to 1.6 uM for FK-3311. Antiinflammatory activity was assessed by inhibition of adjuvantinduced arthritis, and analgesic activity was determined in the acetic acid-induced writhing assay. Following p.o. administration of 10 mg/kg, racemic (2) and its optical isomers showed activity comparable to FK-3311 (76% inhibition) in the adjuvant arthritis test, whereas racemic (5) showed very weak activity, and (R)- and (S)-(5) were not tested. With regard to analgesic effects, FK-3311 and racemic (2) showed 81 and 62% inhibitions, respectively, at a dose of 100 mg/kg p.o. The (R)- and (S)-isomers of (2) and racemic (5) all showed 46% inhibition of writhing syndrome. (R)- and (S)-(5) were less active showing 16 and 20% inhibitions, respectively[1].

in vivo: L-PVR, CO, PaO(2), and WDR were significantly (P



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