

# Vicriviroc (maleate)

## **Catalog No: tcsc1019**

**Available Sizes** 

#### Size: 5mg

Specifications

#### CAS No:

599179-03-0

#### Formula:

 $C_{32}H_{42}F_{3}N_{5}O_{6}$ 

#### Pathway:

Immunology/Inflammation;GPCR/G Protein;Anti-infection

#### **Target:**

CCR;CCR;HIV

#### **Purity / Grade:**

>98%

#### Solubility:

H2O : 25 mg/mL (38.48 mM; Need ultrasonic and warming); DMSO : 50 mg/mL (76.96 mM; Need ultrasonic)

#### **Alternative Names:**

SCH-417690 (maleate);SCH-D (maleate)

#### **Observed Molecular Weight:**

649.7

### **Product Description**

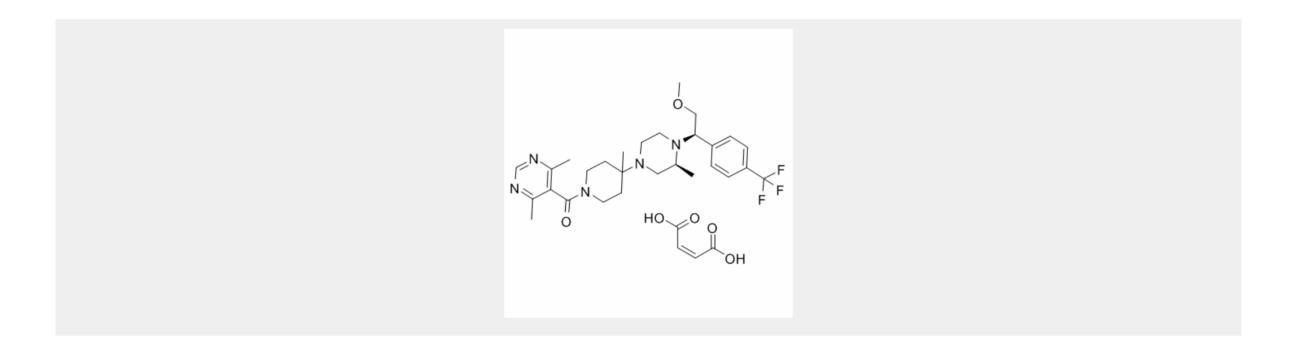
Vicriviroc maleate is a potent, selective, oral bioavailable and CNS penetrated antagonist of **CCR5**, with a  $K_i$  of 2.5 nM, and also inhibits HIV-1 in PBMC cells, with  $IC_{90}$ s of 3.3 nM (JrFL), 2.8 nM (ADA-M), 1.8 nM (301657), 4.9 nM (JV1083) and 10 nM (RU 570). IC50 & Target: Ki: 2.5 nM (CCR5)<sup>[1]</sup>



IC90: 3.3 nM (HIV-1 JrFL, in PBMC cells), 2.8 nM (HIV-1 ADA-M, in PBMC cells), 1.8 nM (HIV-1 301657, in PBMC cells), 4.9 nM (HIV-1 JV1083, in PBMC cells) and 10 nM (HIV-1 RU 570, in PBMC cells)<sup>[1]</sup>

*In Vitro:* Vicriviroc (Sch-417690/Sch-D) is a potent, selective and oral bioavailable inhibitor of CCR5, with a K<sub>i</sub> of 2.5 nM, and also inhibits HIV-1 in PBMC cells, with IC<sub>90</sub>s of 3.3 (JrFL), 2.8 (ADA-M), 1.8 (301657), 4.9 (JV1083) and 10 nM (RU 570). In addition, Vicriviroc shows a mean IC<sub>50</sub> and IC<sub>90</sub> of 0.45 nM and 4 nM for a panel of HIV isolates, and has weak activity against hERG activity (IC  $_{50}$ , 5.8  $\mu$ M)<sup>[1]</sup>. Vicriviroc inhibits chemotactic response to MIP-1 $\alpha$  with IC<sub>50</sub> values below 1 nM, and suppresses RANTES-induced signaling with a mean IC<sub>50</sub> of 4.2 ± 1.3 nM. Vicriviroc potently suppresses all the viral isolates tested, with geometric mean EC<sub>50</sub>s of 0.04-2.3 nM and IC<sub>90</sub>s of 0.45-18 nM<sup>[2]</sup>.

*In Vivo:* Vicriviroc (10 mg/kg) has good oral availablity in rats and monkeys, with no acute CNS or GI effects in rats<sup>[1]</sup>.



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