



Irinotecan (hydrochloride trihydrate)

Catalog No: tcsc1139

3

Available Sizes

Size: 50mg

Size: 100mg

Size: 200mg

Size: 500mg



Specifications

CAS No:

136572-09-3

Formula:

 $C_{33}H_{45}CIN_{4}O_{9}$

Pathway:

Cell Cycle/DNA Damage; Autophagy

Target:

Topoisomerase; Autophagy

Purity / Grade:

>98%

Solubility:

H2O: 1.52 mg/mL (2.24 mM; Need ultrasonic and warming); DMSO: 50 mg/mL (73.84 mM; Need ultrasonic)

Observed Molecular Weight:

677.18

Product Description

Irinotecan hydrochloride trihydrate is a water soluble **topoisomerase I** inhibitor with antitumor activity.





IC50 & Target: Topoisomerase I^[1]

In Vitro: Irinotecan hydrochloride trihydrate is a topoisomerase I inhibitor. Irinotecan inhibits the growth of LoVo and HT-29 cells, with IC $_{50}$ s of 15.8 \pm 5.1 and 5.17 \pm 1.4 μ M, respectively, and induces similar amounts of cleavable complexes in both in LoVo and HT-29 cells^[2]. Irinotecan suppresses the proliferation of human umbilical vein endothelial cells (HUVEC), with an IC $_{50}$ of 1.3 μ M^[3].

In Vivo: Irinotecan (CPT-11, 5 mg/kg) significantly inhibits the growth of tumors by intratumoral injection daily for 5 days, on two consecutive weeks in rats, and such effects also occur via continuous intraperitoneal infusion by osmotic minipump into mice. However, Irinotecan (10 mg/kg) shows no effect on the growth of tumor by i.p^[1]. Irinotecan (CPT-11, 100-300 mg/kg, i.p.) apparently suppresses tumor growth of HT-29 xenografts in athymic female mice by day 21. The two groups of Irinotecan (125 mg/kg) plus TSP-1 (10 mg/kg per day) or Irinotecan (150 mg/kg) in combination TSP-1 (20 mg/kg per day) are nearly equally effective and inhibit tumor growth 84% and 89%, respectively, and both are more effective than Irinotecan alone at doses of 250 and 300 mg/kg^[3].

$$H-CI$$
 H_2O H_2O H_2O

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