



# **Topotecan (Hydrochloride)**

Catalog No: tcsc1498



## **Available Sizes**

Size: 10mg

Size: 50mg

Size: 100mg



# **Specifications**

#### CAS No:

119413-54-6

#### Formula:

 $C_{23}H_{24}CIN_3O_5$ 

## **Pathway:**

Cell Cycle/DNA Damage; Autophagy

#### **Target:**

Topoisomerase; Autophagy

## **Purity / Grade:**

>98%

## **Solubility:**

DMSO: 15.3 mg/mL (33.41 mM; Need ultrasonic and warming)

#### **Alternative Names:**

SKF 104864A;NSC 609669

## **Observed Molecular Weight:**

457.91

# **Product Description**





Topotecan Hydrochloride (SKF 104864A; NSC 609669) is a **Topoisomerase I** inhibitor with potent antineoplastic activities.

IC50 & Target: Topoisomerase I<sup>[1]</sup>

In Vitro: Topotecan (SKF 104864A) obviously inhibits proliferation of not only human glioma cells but also glioma stem cells (GSCs) in a dose- and time-dependent manner. According to the IC $_{50}$  values at 24 h, 3  $\mu$ M of Topotecan (SKF 104864A) is selected as the optimal administration concentration. In addition, Topotecan (SKF 104864A) induces cell cycle arrest in G0/G1 and S phases and promoted apoptosis. Results show that the cell viability is inhibited by Topotecan (SKF 104864A) in a dose-dependent manner. 2, 20 and 40  $\mu$ M of Topotecan (SKF 104864A) obviously inhibits the cell viability compared with the control groups. The IC $_{50}$  values of Topotecan (SKF 104864A) at 24 h are 2.73±0.25  $\mu$ M of U251 cells, 2.95±0.23  $\mu$ M of U87 cells, 5.46±0.41  $\mu$ M of GSCs-U251 and 5.95±0.24  $\mu$ M of GSCs-U87. Thus 3  $\mu$ M of Topotecan (SKF 104864A) is selected as the optimal administration concentration in the subsequent experiments<sup>[1]</sup>.

*In Vivo:* NUB-7 metastatic model, the animals belonging to all the 4 groups are sacrificed after 14 days treatment. Compared with the control, Low dose metronomic (LDM) Topotecan (SKF 104864A) and TP+Pazopanib (PZ) liver weights are significantly lower in TP+PZ-treated animals, compared with PZ. Microscopic tumors are visible in the livers of mice belonging to all the groups except TP+PZ confirming the ability of Topotecan (SKF 104864A)+PZ to control liver metastasis. In a previous dose-response study, the daily dose of oral metronomic Topotecan (SKF 104864A) (0.5, 1.0, and 1.5 mg/kg) causes greater reduction in microvascular density compared with weekly maximum-tolerated dose regimen (7.5 and 15 mg/kg) in an ovarian cancer model, but the mice treated with 1.5 mg/kg daily, oral Topotecan (SKF 104864A) show decreased food intake, and a lesser antitumor effect<sup>[2]</sup>.

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