

# (+) -Apogossypol

**Catalog No: tcsc3563** 

Available Sizes

Size: 5mg

Size: 10mg

**Specifications** 

CAS No:

66389-74-0

#### Formula:

 $C_{28}H_{30}O_6S$ 

#### Pathway:

Apoptosis

#### **Target:**

Bcl-2 Family

#### Purity / Grade:

>98%

**Solubility:** 10 mM in DMSO

### Alternative Names: Apogossypol;NSC736630

**Observed Molecular Weight:** 

494.6

## **Product Description**

(+)-Apogossypol is a **pan-BCL-2** antagonist. (+)-Apogossypol binds to **McI-1**, **BcI-2** and **BcI-xL** with **EC**<sub>50</sub>s of 2.6, 2.8 and 3.69 μM, respectively.

Copyright 2021 Taiclone Biotech Corp.



IC50 & Target: EC50: 2.60 μM (Mcl-1), 2.80 μM (Bcl-2), 3.69 μM (Bcl-xL)<sup>[1]</sup>

Kd: 1.70 μM (Bcl-xL)<sup>[1]</sup>

*In Vitro:* In agreement with NMR binding and fluorescence polarization assays (FPAs) data, (+)-Apogossypol displays potent binding affinity to Bcl-xL with  $K_d$  values of 1.7  $\mu$ M<sup>[1]</sup>.To investigate the inhibitory effects of (+)-Apogossypol and Gossypol on LNCaP cell survival, the MTT assay is performed. The results demonstrate that (+)-Apogossypol inhibits the proliferation of LNCaP cells in a time-and dose-dependent manner, in a similar way with Gossypol. The concentration for 50% inhibition (IC<sub>50</sub>) on LNCaP cells within ~72 h is 9.57  $\mu$ M, while the IC<sub>50</sub> of Gossypol on LNCaP cells is 10.35  $\mu$ M<sup>[2]</sup>.

*In Vivo:* Due to its modified structure, (+)-Apogossypol is expected to exhibit lower toxicity while maintaining the significant antigrowth and anti-tumor activities in vitro, similar to those of Gossypol. The anti-cancer effect of (+)-Apogossypol is evaluated in mice bearing subcutaneous LNCaP cell xenografts. The tumor growth is monitored and measured by a caliper and balance. The survival rate of the mice is notably improved by (+)-Apogossypol. Of note, the tumor sizes are also markedly decreased by (+)-Apogossypol treatment (P[2].



Copyright 2021 Taiclone Biotech Corp.