



YK-4-279

Catalog No: tcsc0667



## **Available Sizes**

Size: 10mg

Size: 50mg

Size: 100mg



## **Specifications**

CAS No:

1037184-44-3

Formula:

 $C_{17}^{H}_{13}^{Cl}_{2}^{NO}_{4}$ 

**Pathway:** 

Cell Cycle/DNA Damage

**Target:** 

**DNA/RNA Synthesis** 

**Purity / Grade:** 

>98%

**Solubility:** 

DMSO: 25 mg/mL (68.27 mM; Need ultrasonic)

**Observed Molecular Weight:** 

366.2

## **Product Description**

YK 4-279 is an inhibitor of RNA Helicase A (RHA) binding to the oncogenic transciption factor EWS-FLI1. YK-4-279 inhibits Ewing\'s sarcoma family tumor (ESFT) cell growth; YK-4-279 induces apoptosis.





IC50 value:

Target: RNA Helicase A

ES-FLI1 is an oncogenic fusion protein found in Ewing's sarcoma, a family of undifferentiated tumors that occur throughout the body. The binding of RNA helicase A (RHA) to ES-FLI1 promotes its oncogenic function. YK-4-279 is an inhibitor of protein-protein interactions between ES-FLI1 and RHA. At 10  $\mu$ M, YK-4-279 blocks RHA binding to ES-FLI1 and induces apoptosis of a panel of Ewing's sarcoma tumor cell lines with IC50 values ranging from 0.5-2  $\mu$ M. At 1.5 mg per dose, YK-4-279 reduces the growth of Ewing's sarcoma orthotopic xenografts in mice after treatment with the inhibitor for two weeks.

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