

# Taltobulin

Catalog No: tcsc3299

Available Sizes

**Specifications** 

Size: 5mg

Size: 10mg

Size: 50mg

Ξ

CAS No:

228266-40-8

Formula:

 $\mathsf{C}_{27}\mathsf{H}_{43}\mathsf{N}_{3}\mathsf{O}_{4}$ 

Pathway:

Cell Cycle/DNA Damage;Cytoskeleton;Antibody-drug Conjugate/ADC Related

# **Target:**

Microtubule/Tubulin;Microtubule/Tubulin;ADC Cytotoxin

## Purity / Grade:

**Solubility:** DMSO : ≥ 100 mg/mL (211.13 mM)

#### **Alternative Names:**

HTI-286;SPA-110

### **Observed Molecular Weight:**

473.65

# **Product Description**

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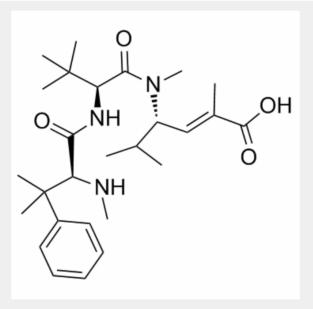
Taltobulin (HTI-286; SPA-110) is an analogue of Hemiasterlin; potent tubulin inhibitor; ADCs cytotoxin.

IC50 value:

Target: tubulin

in vitro: HTI-286 significantly inhibited proliferation of all three hepatic tumor cell lines (mean IC50 = 2 nmol/L +/- 1 nmol/L) in vitro. Interestingly, no decrease in viable primary human hepatocytes (PHH) was detected under HTI-286 exposure [1]. In all cell lines tested, HTI-286 was a potent inhibitor of proliferation and induced marked increases in apoptosis. Despite similar transcriptomic changes regarding cell death and cell cycle regulating genes after exposure to HTI-286 or docetaxel, array analysis revealed distinct molecular signatures for both compounds [2].

in vivo: Intravenous administration of HTI-286 significantly inhibited tumor growth in vivo (rat allograft model) [1]. HTI-286 significantly inhibited growth of PC-3 and LNCaP xenografts and retained potency in PC-3dR tumors. Simultaneous castration plus HTI-286 therapy was superior to sequential treatment in the LNCaP model [2].



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