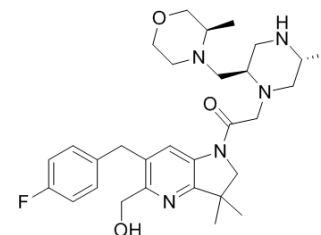


## ASTX660

<b>Cat. No.:</b>	HY-109565		
<b>CAS No.:</b>	1799328-86-1		
<b>Molecular Formula:</b>	C <sub>30</sub> H <sub>42</sub> FN <sub>5</sub> O <sub>3</sub>		
<b>Molecular Weight:</b>	539.68		
<b>Target:</b>	IAP		
<b>Pathway:</b>	Apoptosis		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : ≥ 50 mg/mL (92.65 mM)  
 \* "≥" means soluble, but saturation unknown.

Concentration	Mass		
	1 mg	5 mg	10 mg
<b>1 mM</b>	1.8529 mL	9.2647 mL	18.5295 mL
<b>5 mM</b>	0.3706 mL	1.8529 mL	3.7059 mL
<b>10 mM</b>	0.1853 mL	0.9265 mL	1.8529 mL

Please refer to the solubility information to select the appropriate solvent.

#### In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
 Solubility: ≥ 2.5 mg/mL (4.63 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
 Solubility: ≥ 2.5 mg/mL (4.63 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
 Solubility: ≥ 2.5 mg/mL (4.63 mM); Clear solution

### BIOLOGICAL ACTIVITY

#### Description

ASTX660 is an orally bioavailable dual antagonist of cellular inhibitor of apoptosis protein (cIAP) and X-linked inhibitor of apoptosis protein (XIAP).

#### IC<sub>50</sub> & Target

cIAP, XIAP<sup>[1]</sup>

#### In Vitro

ASTX660 is an orally bioavailable dual antagonist of cIAP and XIAP, currently being investigated in a single-agent Phase 1/2

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clinical trial in patients with advanced solid tumors and lymphomas. Twenty-one triple-negative breast cancer (TNBC) cell lines are treated with ASTX660 in vitro and it is found that 43% are sensitive to ASTX660<sup>[1]</sup>.  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

**In Vivo**

In HCC1806 xenografts in mice, ASTX660 (daily oral treatment) causes moderate tumor growth inhibition but not regression [1].  
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## CUSTOMER VALIDATION

- Cell Reports Medicine. 2020 Jun 23;1(3):100037.

See more customer validations on [www.MedChemExpress.com](http://www.MedChemExpress.com)

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## REFERENCES

[1]. Tomoko Smyth, et al. Abstract 1287: The dual IAP antagonist, ASTX660, increases the anti-tumor activity of NSC 125973 in preclinical models of triple-negative breast cancer in vivo. Cancer Res 2016;76(14 Suppl).

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**Caution: Product has not been fully validated for medical applications. For research use only.**

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