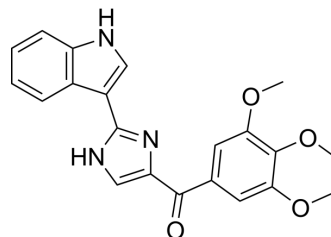


## Sabizabulin

<b>Cat. No.:</b>	HY-120599
<b>CAS No.:</b>	1332881-26-1
<b>Molecular Formula:</b>	C <sub>21</sub> H <sub>19</sub> N <sub>3</sub> O <sub>4</sub>
<b>Molecular Weight:</b>	377.39
<b>Target:</b>	Microtubule/Tubulin; Apoptosis; HPV
<b>Pathway:</b>	Cell Cycle/DNA Damage; Cytoskeleton; Apoptosis; Anti-infection
<b>Storage:</b>	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)



### SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 20 mg/mL (53.00 mM; Need ultrasonic)					
		Solvent Concentration	Mass			
	<b>Preparing Stock Solutions</b>			1 mg	5 mg	10 mg
		1 mM		2.6498 mL	13.2489 mL	26.4978 mL
		5 mM		0.5300 mL	2.6498 mL	5.2996 mL
	10 mM		0.2650 mL	1.3249 mL	2.6498 mL	
Please refer to the solubility information to select the appropriate solvent.						
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2 mg/mL (5.30 mM); Clear solution  2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2 mg/mL (5.30 mM); Suspended solution; Need ultrasonic					

### BIOLOGICAL ACTIVITY

<b>Description</b>	VERU-111 (ABI-231) is a potent and orally active $\alpha$ and $\beta$ tubulin inhibitor, which displays strong antiproliferative activity, with an average IC <sub>50</sub> of 5.2 nM against panels of melanoma and prostate cancer cell lines. VERU-111 (ABI-231) suppresses tumor growth and metastatic phenotypes of cervical cancer cells via targeting HPV E6 and E7, and has potential for the treatment of prostate cancer <sup>[1][2][3]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	tubulin <sup>[1]</sup>
<b>In Vitro</b>	VERU-111 (2.5-80 nM; 24-48 hours) inhibits Panc-1, AsPC-1 and HPAF-II cells growth in a dose and time-dependent manner (24 hours: IC <sub>50</sub> s of 25, 35 and 35?nM, respectively; 48 hours: IC <sub>50</sub> s of 11.8, 15.5, and 25?nM, respectively) <sup>[4]</sup> . ?VERU-111 (5-20 nM; 24 hours) arrests Panc-1 and AsPC-1 cells in G2/M phase in a dose-dependent manner <sup>[4]</sup> . ?VERU-111 (5-20?nM; 24 hours) shows dose-dependent inhibition of pro-Caspase 3 and 9 and activation of Caspase-3 and 9,

induces the expression of Bax and Bad, and inhibits the expression of Bcl-2 and Bcl-xl proteins in both AsPC-1 and Panc-1 cells<sup>[4]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### Cell Proliferation Assay<sup>[4]</sup>

Cell Line:	Panc-1, AsPC-1, HPAF-II cells
Concentration:	2.5, 5, 10, 20, 40, 80 nM
Incubation Time:	24, 48 hours
Result:	Inhibited the growth of PanCa cells in a dose and time-dependent manner. The IC <sub>50</sub> of VERU-111 was 25, 35 and 35 nM in Panc-1, AsPC-1 and HPAF-II, respectively after 24 h treatment, while 48 h post-treatment it was 11.8, 15.5, and 25 nM.

#### Apoptosis Analysis<sup>[4]</sup>

Cell Line:	Panc-1, AsPC-1 cells
Concentration:	5, 10, 20 nM
Incubation Time:	24 hours
Result:	Arrested Panc-1 and AsPC-1 cells in G2/M phase in a dose-dependent manner.

#### Western Blot Analysis<sup>[4]</sup>

Cell Line:	AsPC-1 and Panc-1 cells
Concentration:	5, 10, 20 nM
Incubation Time:	24 hours
Result:	Dose-dependent inhibition of pro-Caspase 3 and 9 and activation of Caspase-3 and 9 in both AsPC-1 and Panc-1 cells. Induces the expression of Bax and Bad and inhibited the expression of Bcl-2 and Bcl-xl proteins.

#### In Vivo

VERU-111 (50 µg/mouse; intra-tumorally; 3 times per week for 3 weeks) effectively inhibits tumor growth as compared to vehicle-treated group. None of the mouse showed any apparent toxicity as constant increase of body weight in VERU-111 treated mice<sup>[4]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Six-week-old female athymic nude mice (bearing AsPC-1 cells)
Dosage:	50 µg/mouse
Administration:	Intra-tumorally; 3 times per week for 3 weeks
Result:	Effectively inhibited tumor growth.

#### CUSTOMER VALIDATION

- bioRxiv. 2023 Jun 30.

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## REFERENCES

- [1]. Wang Q, et al. Structure-Guided Design, Synthesis, and Biological Evaluation of (2-(1H-Indol-3-yl)-1H-imidazol-4-yl)(3,4,5-trimethoxyphenyl) Methanone (ABI-231) Analogues Targeting the Colchicine Binding Site in Tubulin. *J Med Chem.* 2019 Jul 12.
- [2]. Qinghui Wang, et al. Discovery of ABI-231 analogs targeting the colchicine site in tubulin for advanced melanoma. *Cancer Research* 76(14 Supplement):4848-4848.
- [3]. Vivek Kashyap, et al. ABI-231: A novel small molecule suppresses tumor growth and metastatic phenotypes of cervical cancer cells via targeting Human papilloma virus (HPV) E6 and E7. *Cancer Research* 78(13 Supplement):679-679.
- [4]. Kashyap VK, et al. Therapeutic efficacy of a novel  $\beta$ III/ $\beta$ IV-tubulin inhibitor (VERU-111) in pancreatic cancer. *J Exp Clin Cancer Res.* 2019 Jan 23;38(1):29.
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**Caution: Product has not been fully validated for medical applications. For research use only.**

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