# **Product** Data Sheet

### **TACC3** inhibitor 1

Cat. No.:HY-151985Molecular Formula: $C_{20}H_{21}N_5OS$ Molecular Weight:379.48

Target: Apoptosis
Pathway: Apoptosis

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

#### **BIOLOGICAL ACTIVITY**

Description

TACC3 inhibitor 1 is a potent and cross the blood-brain barrier TACC3 inhibitor. TACC3 inhibitor 1 induces Apoptosis and cell cycle arrest at G2/M phase. TACC3 inhibitor 1 induces the generation of intracellular ROS. TACC3 inhibitor 1 shows antiproliferative and anti-tumor activity<sup>[1]</sup>.

In Vitro

TACC3 inhibitor 1 (compound 7g) (1, 2, 4  $\mu$ M; 36 h) induces apoptosis and cell cycle arrest at G2/M phase<sup>[1]</sup>. TACC3 inhibitor 1 (1, 2, 4  $\mu$ M; 36 h) decreases the mitochondrial membrane potential and induces the generation of intracellular ROS<sup>[1]</sup>.

TACC3 inhibitor 1 (1, 2, 4  $\mu$ M; 24 h) inhibits cell migration and invasion in U87 cells<sup>[1]</sup>.

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

Cell Proliferation  $Assay^{[1]}$ 

Cell Line:	U251, U87, MDA-MB-231, JIMT-1, SKOV-3, HLF-1 cells	
Concentration:	0-100 μΜ	
Incubation Time:	24 h	
Result:	Showed antiproliferative activities with IC $_{50}$ s of 5.61, 3.29, 6.46, 4.84, 8.21, 23.11 $\mu$ M for U251, U87, MDA-MB-231, JIMT-1, SKOV-3, HLF-1 cells, respectively.	
Cell Cycle Analysis <sup>[1]</sup>		

Cell Line:	U87 cells	
Concentration:	1, 2, 4 μΜ	
Incubation Time:	36 h	
Result:	Induced cell cycle arrest at G2/M phase with the 41.2% cells at G2/M phase at 4 $\mu$ M.	

## Apoptosis Analysis<sup>[1]</sup>

Cell Line:	U87 cells
Concentration:	1, 2, 4 μΜ

	Incubation Time:	36 h	
	Result:	Induced apoptosis in a dose-dependent manner with the percentage of early apoptotic cells increased from 4.49% in control group to 21.42%, late apoptotic cells increased from 0.52% to 26.72% at 4 $\mu$ M.	
In Vivo	TACC3 inhibitor 1 (20 mg/kg; i.p.; daily for 20 days) shows anti-tumor activity in mouse <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
	Animal Model:	Six-week-old BALB/c nude mice (U87 xenograft model) <sup>[1]</sup>	
	Dosage:	20 mg/kg	
	Administration:	I.p.; daily for 20 days	
	Result:	Exhibited a significant tumor growth regression and no observable toxicity during the administration period and had no effects on the body weight.	

#### **REFERENCES**

[1]. Zhao W, et al. Discovery of novel analogs of KHS101 as transforming acidic coiled coil containing protein 3 (TACC3) inhibitors for the treatment of glioblastoma. Eur J Med Chem. 2022 Dec 15;244:114874.

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

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