



SZABO SCANDIC

Part of Europa Biosite

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!
See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

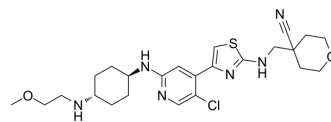
mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

JSH-150

Cat. No.:	HY-X0150		
CAS No.:	2247481-21-4		
Molecular Formula:	C ₂₄ H ₃₃ ClN ₆ O ₂ S		
Molecular Weight:	505.08		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (197.99 mM; Need ultrasonic)						
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg	
				1 mM	1.9799 mL	9.8994 mL	19.7988 mL
				5 mM	0.3960 mL	1.9799 mL	3.9598 mL
				10 mM	0.1980 mL	0.9899 mL	1.9799 mL
Please refer to the solubility information to select the appropriate solvent.							
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (4.95 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (4.95 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (4.95 mM); Clear solution						

BIOLOGICAL ACTIVITY

Description	JSH-150 is a highly selective and potent CDK9 inhibitor with an IC ₅₀ of 1 nM.
IC ₅₀ & Target	IC ₅₀ : 1 nM (CDK9/Cyclin T1), 292 nM (CDK16/Cyclin Y), 1.34 μM (CDK1/Cyclin B), 1.68 μM (CDK14/Cyclin Y), 1.72 μM (CDK7/Cyclin H/MNAT1), 2.86 μM (CDK2/Cyclin A), 4.64 μM (CDK5/p25) ^[1]
In Vitro	The antiproliferative effect of JSH-150 is examined on a panel of established cancer cell lines. JSH-150 exhibits potent antiproliferative activities in solid tumor cell lines such as A375 (melanoma), A431 (squamous), BE(2)M17 (neuroblastoma), GIST-T1 (GIST) and COLO205 (colon cancer) with GI ₅₀ values from 0.002 to 0.044 μM. In the leukemia cell lines including acute myeloid leukemia (AML), chronic lymphocytic leukemia (CLL) and B cell lymphoma cell lines, JSH-150 also displays

strong growth inhibition efficacies with GI₅₀s ranging from single to double digit nM. In addition, JSH-150 is much less sensitive in normal CHO cells (GI₅₀: 1.1 μM) compared with the cancer cell lines^[1].

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

In Vivo

Treatment of JSH-150 at all dosages, i.e., 10, 20 and 30 mg/kg/day, can almost completely suppress the tumor progression in the first two weeks and does not affect the animal's weight indicating that there is no general cytotoxicity at these doses. After stopping the treatment of JSH-150, the tumors of the animals treated with 10 mpk drug dosage start to grow again. However, this tumor recurrence is not observed in the 20 and 30 mpk dosage groups during the following week after administration of JSH-150 is stopped and p values are quantified on the 21st day, which are 0.042, 0.0035 and 0.0028, respectively. The PK properties of JSH-150 are evaluated in different species including mice, Sprague-Dawley rats and beagle dogs through intravenous injection and oral administration. JSH-150 is absorbed rapidly in dogs and mice (T_{max} =1.33 h and 2.00 h respectively) but slowly in rats (T_{max}=3.33 h). JSH-150 also displays different half-lives in three different species via oral administration (T_{1/2}=1.55 h in mice, 3.37 h in rats and 20.37 h in dogs), which indicates that it is metabolized very slowly in dogs compared with mice and rats. In addition, JSH-150 exhibits acceptable bioavailability in mice, rats and dogs (F=45.01%, 45.10% and 39.15%, respectively). The PK properties indicated that JSH-150 is suitable for oral administration^[1].

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

PROTOCOL

Cell Assay ^[1]

The A375 (melanoma), A431 (squamous), BE(2)M17 (neuroblastoma), BE(2)M17 (neuroblastoma), CRL-2234 (hepatoma), COLO205 (colon cancer), A549 (lung adenocarcinoma), Ramos (B cell lymphoma), MV4-11 (AML), Ramos (B cell lymphoma), U937 (AML), CHL (hamster lung cell), and CHO (hamster ovary cell) cell lines are used. OCI-AML-3 (AML), SKM-1 (AML), MEC-1 (CLL), MEC-2 (CLL) and HL-60 (human promyelocytic leukemia cells) are used. Human GIST-T1 cells are used. MOLM-13 and MOLM14 cell lines are used. All the cells are grown in a humidified incubator at 37°C under 5% CO₂. A375, A431, GIST-T1, A549, Colo205 and CHO cells are maintained in DMEM supplemented with 10% FBS, 1% Penicillin/Streptomycin. BE(2)M17 cells are cultured with 1:1 mixture of ATCC-formulated Eagle's minimum essential medium, and F12 Medium. MV4-11, MEC-1 and MEC-2 are grown in IMDM supplemented with 10% FBS, 1% Penicillin/Streptomycin. CRL-2234, U2932, U937, Ramos, MOLM13, MOLM14, OCI-AML-3, SKM-1, HL-60 and CHL are grown in RPMI 1640 medium supported with 10% FBS and 1% Penicillin/Streptomycin. Adherent cells are grown in tissue culture flasks until they are 85-95% confluent prior to use. For suspension cells, cells are collected by spin down at 800 rpm/min for 5 min before use. Cells are grown in 96-well culture plates (3000 cells/well). The compounds (e.g., JSH-150) at various concentrations are added into the plates. Cell proliferation is determined after treatment with compounds (e.g., JSH-150) for 72 h. Cell viability is measured using the Cell Titer-Glo assay and luminescence is measured in a multilabel reader^[1].

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

Animal Administration ^[1]

Mice^[1]

Five-week-old female nu/nu mice are used. Prior to implantation, cells are harvested during exponential growth. Five million MV4-11 cells in PBS are formulated as a 1:1 mixture with Matrigel and injected into the subcutaneous space on the right flank of nu/nu mice. Daily oral administration is initiated when MV4-11 tumors have reached a size of 200-400 mm³. Animals are then randomized into treatment groups of 5 mice each for efficacy studies. JSH-150 is delivered daily in a HKI solution (0.5% methocellulose/0.4% Tween80 in ddH₂O) by oral gavage. A range of doses of JSH-150 or its vehicle as control are administered. Female nu/nu mice bearing established MV4-11 tumor xenografts are treated with JSH-150 at 10, 20, and 30 mg/kg/d dosage or vehicle. Body weight is measured daily and tumor growth is measured every day after JSH-150 treatment. Tumor volume is calculated^[1].

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

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA