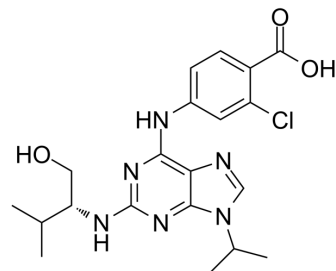


Purvalanol B

Cat. No.:	HY-18299		
CAS No.:	212844-54-7		
Molecular Formula:	C ₂₀ H ₂₅ ClN ₆ O ₃		
Molecular Weight:	432.9		
Target:	CDK; Parasite		
Pathway:	Cell Cycle/DNA Damage; Anti-infection		
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 : ≥ 40 mg/mL (92.40 mM)
 * "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent		1 mg	5 mg	10 mg
	Concentration	Mass			
	1 mM		2.3100 mL	11.5500 mL	23.1000 mL
	5 mM		0.4620 mL	2.3100 mL	4.6200 mL
	10 mM		0.2310 mL	1.1550 mL	2.3100 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 (5.78 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
 Solubility: ≥ 2.5 mg/mL (5.78 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

Purvalanol B (NG 95) is a potent, selective, reversible and ATP-competitive inhibitor CDK, with IC₅₀s of 6 nM, 6 nM, 9 nM, 6 nM for cdc2-cyclin B, CDK2-cyclin A, CDK2-cyclin E and CDK5-p35, respectively. Purvalanol B shows selectivity for CDK over a range of other protein kinases (IC₅₀>1000 nM). Purvalanol B inhibits the growth a chloroquine-resistant strain of *P. falciparum*^{[1][1]}.

IC₅₀ & Target

cdc2/cyclin B 6 nM (IC ₅₀)	cdk2/cyclin A 6 nM (IC ₅₀)	CDK2/cyclinE 9 nM (IC ₅₀)	CDK5/p35 6 nM (IC ₅₀)
-------------------------------------------	-------------------------------------------	------------------------------------------	--------------------------------------

In Vitro

Purvalanol B binds to *P. falciparum* casein kinase 1 (CK1) from blood stage cell lysates and inhibits the growth a

chloroquine-resistant strain of *P. falciparum* (FCR-3) with an IC₅₀ of 7.07 μM^[1].

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

CUSTOMER VALIDATION

- Anticancer Drugs. 2022 Aug 9.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Gray NS, et, al. Exploiting chemical libraries, structure, and genomics in the search for kinase inhibitors. *Science*. 1998 Jul 24;281(5376):533-8.

[2]. Bullard KM, et, al. Effects of cyclin-dependent kinase inhibitor Purvalanol B application on protein expression and developmental progression in intra-erythrocytic *Plasmodium falciparum* parasites. *Malar J*. 2015 Apr 8;14:147.

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