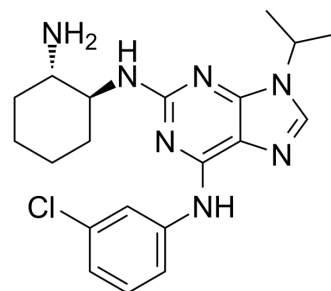


SRI-29329

Cat. No.:	HY-123600		
CAS No.:	2086809-58-5		
Molecular Formula:	C ₂₀ H ₂₆ ClN ₇		
Molecular Weight:	399.92		
Target:	CDK		
Pathway:	Cell Cycle/DNA Damage		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 50 mg/mL (125.03 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
			1 mM	2.5005 mL	12.5025 mL	25.0050 mL
			5 mM	0.5001 mL	2.5005 mL	5.0010 mL
			10 mM	0.2501 mL	1.2502 mL	2.5005 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 (6.25 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (6.25 mM); Suspended solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description	SRI-29329 is a specific CLK inhibitor, with IC ₅₀ values of 78 nM, 16 nM and 86 nM for CLK1, CLK2 and CLK4, respectively ^[1] .
In Vitro	SRI-29329 (compound 8) shows some modest (~5 fold) selectivity for CLK2 over CLK1 and CLK4 without significant CDK1,4,6 activity ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Yihui Shi, et al. A triple exon-skipping luciferase reporter assay identifies a new CLK inhibitor pharmacophore. *Bioorg Med Chem Lett.* 2017 Feb 1;27(3):406-412.

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

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