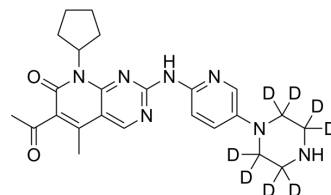


Palbociclib-d₈

Cat. No.:	HY-50767S		
CAS No.:	1628752-83-9		
Molecular Formula:	C ₂₄ H ₂₁ D ₈ N ₇ O ₂		
Molecular Weight:	455.58		
Target:	CDK		
Pathway:	Cell Cycle/DNA Damage		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro

H₂O : 8.33 mg/mL (18.28 mM; ultrasonic and warming and adjust pH to 3 with HCl and heat to 80°C)

DMSO : 1 mg/mL (2.20 mM; ultrasonic and warming and heat to 60°C)

DMSO : 1 mg/mL (2.20 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent	1 mg	5 mg	10 mg
	Concentration	Mass		
1 mM		2.1950 mL	10.9750 mL	21.9500 mL
5 mM		0.4390 mL	2.1950 mL	4.3900 mL
10 mM		0.2195 mL	1.0975 mL	2.1950 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description

Palbociclib-d₈ is a deuterium labeled Palbociclib. Palbociclib is a selective and orally active CDK4 and CDK6 inhibitor with IC₅₀s of 11 and 16 nM, respectively. Palbociclib has the potential for ER-positive and HER2-negative breast cancer research[1].

IC₅₀ & Target

Cdk4/cyclin D3 9 nM (IC ₅₀)	Cdk4/cyclin D1 11 nM (IC ₅₀)	Cdk6/cyclin D2 16 nM (IC ₅₀)	DYRK1A 2000 nM (IC ₅₀)
MAPK 8000 nM (IC ₅₀)			

REFERENCES

[1]. Fry DW, et al. Specific inhibition of cyclin-dependent kinase 4/6 by PD 0332991 and associated antitumor activity in human tumor xenografts. Mol Cancer Ther. 2004 Nov;3(11):1427-38.

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

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