

Product Name: PD 0332991 (Palbociclib) Revision Date: 01/10/2021



PD 0332991 (Palbociclib)

Cat. No.:	B7798
CAS No.:	571190-30-2
Formula:	C24H29N7O2
M.Wt:	447.53
Synonyms:	PD0332991;PD-0332991;PD 0332991
Target:	Cell Cycle/Checkpoint
Pathway:	Cyclin-Dependent Kinases
Storage:	Store at -20°C

Solvent & Solubility

	insoluble in EtOH; in	soluble in EtOH; insoluble in H2O; insoluble in DMSO					
Preparing In Vitro Stock Solution	Preparing	Mass Solvent Concentration	1mg	5mg	10mg		
	Stock Solutions	1 mM	2.2345 mL	11.1724 mL	22.3449 mL		
		5 mM	0.4469 mL	2.2345 mL	4.4690 mL		
		10 mM	0.2234 mL	1.1172 mL	2.2345 mL		

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

Biological Activity

Shortsummary	CDK4/6 inhibitor, highly selective			
IC ₅₀ & Target	11 nM (CDK4/cyclin D1), 16 nM (CDK6/cyclin D2)			
In Vitro	Cell Viability Assay			
	Cell Line:	MDA-MB-453 cells		
	Preparation method:	The solubility of this compound in DMSO is		
	Reacting conditions:	80 nM, 24 hours		
	Applications:	MDA-MB-453 breast carcinoma cells exposed to varying concentrations of F		
		0332991 for 24 hours showed a significant increase in the percentage of cells		

		in G1 in the presence of as little as 0.04 $\mu mol/L$ PD 0332991 with a concomitant			
		decline in other phases of the cell cycle. Maximum effects were attained at 0.08			
		$\mu mol/L$ and an exclusive G1 arrest was maintained even at concentrations as			
		high as 10 $\mu mol/L,$ consistent with the complete absence of any other effects			
		on the cell cycle.			
	Animal experiment				
In Vivo	Animal models:	Mice bearing Colo-205 colon carcinoma xenografts			
	Dosage form:	Oral administration, 150 or 75 mg/kg, daily for 14 days			
	Applications:	Administration of PD 0332991 produced rapid tumor regressions and a			
		corresponding tumor growth delay of about 50 days with >1 log of tumor cell kill			
		at the highest dose tested. At 37.5 mg/kg, the tumor slowly regressed during			
		treatment. Even at doses as low as 12.5 mg/kg, a 13-day growth delay was			
		obtained indicating a 90% inhibition of tumor growth rate.			
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may			
		slightly differ with the theoretical value. This is caused by an experimental			
		system error and it is normal.			

Product Citations

1. Cingöz O, Goff SP. "Cyclin-dependent kinase activity is required for type linterferon production." Proc Natl Acad Sci U S A. 2018 Mar 27;115(13):E2950-E2959.PMID:29507205

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References

[1] Fry D W, Harvey P J, Keller P R, et al. Specific inhibition of cyclin-dependent kinase 4/6 by PD 0332991 and associated antitumor activity in human tumor xenografts. Molecular cancer therapeutics, 2004, 3(11): 1427-1438.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.



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