Pomalidomide-PEG3-azide

MedChemExpress

®

Cat. No.:	HY-137538	
CAS No.:	2267306-15-8	
Molecular Formula:	$C_{21}H_{24}N_6O_8$	
Molecular Weight:	488.45	
Target:	E3 Ligase Ligand-Linker Conjugates	-N ² N ⁺ N 0000
Pathway:	PROTAC	
Storage:	-20°C, stored under nitrogen, away from moisture	
	* In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen, away from moisture)	

SOLVENT & SOLUBILITY

	Preparing Stock Solutions	Mass Solvent Concentration	1 mg	5 mg	10 mg	
		1 mM	2.0473 mL	10.2365 mL	20.4729 mL	
		5 mM	0.4095 mL	2.0473 mL	4.0946 mL	
		10 mM	0.2047 mL	1.0236 mL	2.0473 mL	
Plea	Please refer to the solubility information to select the appropriate solvent.					
Vivo 1. A	1. Add each solvent one by one: 10% DMSO >> 90% corn oil					
	Solubility: ≥ 5 mg/mL (10.24 mM); Clear solution					

BIOLOGICAL ACTIVITY		
Description	Pomalidomide-PEG3-azide is a synthesized E3 ligase ligand-linker conjugate that incorporates the Pomalidomide based cereblon ligand and 3-unit PEG linker used in PROTAC technology ^[1] . Pomalidomide-PEG3-azide is a click chemistry reagent, it contains an Azide group and can undergo copper-catalyzed azide-alkyne cycloaddition reaction (CuAAc) with molecules containing Alkyne groups. Strain-promoted alkyne-azide cycloaddition (SPAAC) can also occur with molecules containing DBCO or BCN groups.	
IC ₅₀ & Target	Cereblon	
In Vitro	PROTACs contain two different ligands connected by a linker; one is a ligand for an E3 ubiquitin ligase and the other is for the target protein. PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

Product Data Sheet

REFERENCES

[1]. Fangqing Zhang, et al. Discovery of a new class of PROTAC BRD4 degraders based on a dihydroquinazolinone derivative and lenalidomide/pomalidomide. Bioorg Med Chem. 2020 Jan 1;28(1):115228.

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

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