Propargyl-PEG5-amine

MedChemExpress

®

Cat. No.:	HY-126976
CAS No.:	1589522-46-2
Molecular Formula:	C ₁₃ H ₂₅ NO ₅
Molecular Weight:	275.34
Target:	ADC Linker; PROTAC Linkers
Pathway:	Antibody-drug Conjugate/ADC Related; PROTAC
Storage:	-20°C, protect from light * In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

SOLVENT & SOLUBILITY

Prepa	DMSO : 100 mg/mL (363.19 mM; Need ultrasonic)						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	3.6319 mL	18.1594 mL	36.3187 mL		
		5 mM	0.7264 mL	3.6319 mL	7.2637 mL		
		10 mM	0.3632 mL	1.8159 mL	3.6319 mL		
	Please refer to the so	lubility information to select the ap	propriate solvent.				
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (9.08 mM); Clear solution						
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (9.08 mM); Clear solution						
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (9.08 mM); Clear solution						

BIOLOGICAL ACTIVITY				
Description	Propargyl-PEG5-amine is a non-cleavable ADC linker used in the synthesis of antibody-drug conjugates (ADCs). Propargyl- PEG5-amine is a PEG-based PROTAC linker can be used in the synthesis of PROTACs ^[1] . Propargyl-PEG5-amine is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAc) with molecules containing Azide groups.			
IC ₅₀ & Target	Non-cleavable Linker PEGs			
In Vitro	ADCs are comprised of an antibody to which is attached an ADC cytotoxin through an ADC linker.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.			

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Product Data Sheet

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PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins.

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

REFERENCES

[1]. Robert Zamboni, et al. Raf-degrading conjugate compounds. WO2018200981A1.

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

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