

Product Data Sheet

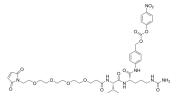
Mal-PEG4-Val-Cit-PAB-PNP

Cat. No.:HY-140145CAS No.:2112738-09-5Molecular Formula: $C_{40}H_{53}N_7O_{15}$ Molecular Weight:871.89Target:ADC Linker

Pathway: Antibody-drug Conjugate/ADC Related

Storage: -20°C, protect from light

* In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)



SOLVENT & SOLUBILITY

In Vitro

DMSO: 100 mg/mL (114.69 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.1469 mL	5.7347 mL	11.4693 mL
	5 mM	0.2294 mL	1.1469 mL	2.2939 mL
	10 mM	0.1147 mL	0.5735 mL	1.1469 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: ≥ 2.08 mg/mL (2.39 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: \geq 2.08 mg/mL (2.39 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	${\it Mal-PEG4-Val-Cit-PAB-PNP is a cleavable 4 unit PEG ADC linker used in the synthesis of antibody-drug conjugates (ADCs)} {\it [1]}.$		
IC ₅₀ & Target	Protease Cleavable	Cleavable	
In Vitro	ADCs are comprised of an antibody to which is attached an ADC cytotoxin through an ADC linker ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		

REFERENCES

1]. Beck A, et al. Strategies and	challenges for the next generatio	n of antibody-drug conjugates. I	Nat Rev Drug Discov. 2017 May;16(5):	315-337.
			al applications. For research use	
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