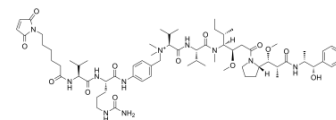


MC-Val-Cit-PAB-Auristatin E

Cat. No.:	HY-128899
CAS No.:	2055896-77-8
Molecular Formula:	C ₆₈ H ₁₀₈ N ₁₁ O ₁₃
Molecular Weight:	1287.65
Target:	Drug-Linker Conjugates for ADC
Pathway:	Antibody-drug Conjugate/ADC Related
Storage:	4°C, sealed storage, away from moisture and light * The compound is unstable in solutions, freshly prepared is recommended.



SOLVENT & SOLUBILITY

In Vitro	DMSO : 200 mg/mL (155.32 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Concentration	Mass	1 mg	5 mg	10 mg
		1 mM		0.7766 mL	3.8830 mL	7.7661 mL
		5 mM		0.1553 mL	0.7766 mL	1.5532 mL
		10 mM		0.0777 mL	0.3883 mL	0.7766 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: ≥ 5 mg/mL (3.88 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 5 mg/mL (3.88 mM); Clear solution					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5 mg/mL (3.88 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	MC-Val-Cit-PAB-Auristatin E is a drug-linker conjugate for ADC with potent antitumor activity by using Auristatin E (a cytotoxic tubulin modifier), linked via the ADC linker MC-Val-Cit-PAB.
IC₅₀ & Target	Auristatin

REFERENCES

[1]. Staben LR, et al. Targeted drug delivery through the traceless release of tertiary and heteroaryl amines from antibody-drug conjugates. Nat Chem. 2016 Dec;8(12):1112-1119.

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

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