SC239

Cat. No.:	HY-148194
CAS No.:	1977557-97-3
Molecular Formula:	C ₆₅ H ₈₄ N ₁₀ O ₁₁
Molecular Weight:	1181.42
Target:	Drug-Linker Conjugates for ADC
Pathway:	Antibody-drug Conjugate/ADC Related
Storage:	4°C, protect from light
	* In solvent : -80°C, 6 months; -20°C, 1 month (protect from light)

SOLVENT & SOLUBILITY

MedChemExpress

	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	0.8464 mL	4.2322 mL	8.4644 mL
		5 mM	0.1693 mL	0.8464 mL	1.6929 mL
		10 mM	0.0846 mL	0.4232 mL	0.8464 mL

BIOLOGICAL ACTIVITY				
Description	SC239 is a cleavable 2-aminophenyl hemiasterlin agent-linker. SC239 can be as the agent-linker for ADC ^[1] . SC239 is a click chemistry reagent, it contains a DBCO group that can undergo strain-promoted alkyne-azide cycloaddition (SPAAC) with molecules containing Azide groups.			
In Vitro	SC239 is composed of a tubulin-targeting 3-aminophenyl hemiasterlin warhead, <u>SC209</u> (HY-144880), and a cleavable valine citrulline p-aminobenzyl carbamate linker functionalized with dibenzocyclooctyne (DBCO) ^[1] . The SC239 drug-linker is conjugated via a cleavable valine citrulline p-aminobenzyl carbamate linker functionalized with dibenzocyclooctyne (DBCO) ^[1] . SC239 conjugates exhibits better cytotoxic activity on Igrov1 cells which have lower expression levels of FoIRα ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

REFERENCES

[1]. Xiaofan Li, et al. Abstract 1782: Discovery and activity of STRO-002, a novel ADC targeting folate receptor alpha for ovarian and endometrial cancer. Cancer Res (2018) 78 (13_Supplement): 1782.

Product Data Sheet

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

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