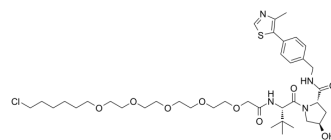


VH032-PEG5-C6-Cl

Cat. No.:	HY-112495
CAS No.:	1799506-06-1
Molecular Formula:	C ₃₈ H ₅₉ ClN ₄ O ₉ S
Molecular Weight:	783.41
Target:	E3 Ligase Ligand-Linker Conjugates
Pathway:	PROTAC
Storage:	-80°C, protect from light, stored under nitrogen



SOLVENT & SOLUBILITY

In Vitro

DMSO : 270 mg/mL (344.65 mM; Need ultrasonic)
Ethanol : 100 mg/mL (127.65 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Concentration	Mass		
		1 mg	5 mg	10 mg
	1 mM	1.2765 mL	6.3824 mL	12.7647 mL
	5 mM	0.2553 mL	1.2765 mL	2.5529 mL
	10 mM	0.1276 mL	0.6382 mL	1.2765 mL

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
Solubility: ≥ 5 mg/mL (6.38 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
Solubility: ≥ 5 mg/mL (6.38 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil
Solubility: ≥ 5 mg/mL (6.38 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

VH032-PEG5-C6-Cl (HaloPROTAC 2) is a conjugate of ligands for E3 and 21-atom-length linker. The connector of linker is Halogen group. VH032-PEG5-C6-Cl incorporates the VH032 based VHL ligand and 5-unit PEG linker. VH032-PEG5-C6-Cl is capable of inducing the degradation of GFP-HaloTag7 in cell-based assays^[1].

IC₅₀ & Target

VHL

In Vitro

Treatment (24 hour) with VH032-PEG5-C6-Cl leads to nearly 70% degradation of GFP-Halotag7 at 2.5 μM^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Buckley DL, et al. HaloPROTACS: Use of Small Molecule PROTACs to Induce Degradation of HaloTag Fusion Proteins. ACS Chem Biol. 2015 Aug 21;10(8):1831-7.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA