

Product Data Sheet

Thalidomide-NH-PEG4-COOH

Cat. No.: HY-134591 CAS No.: 2412056-48-3 Molecular Formula: $C_{24}H_{31}N_3O_{10}$ Molecular Weight: 521.52

Target: E3 Ligase Ligand-Linker Conjugates

Pathway: PROTAC

Storage: -20°C, protect from light

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

SOLVENT & SOLUBILITY

In Vitro

DMSO: 80 mg/mL (153.40 mM; Need ultrasonic) H₂O: 10 mg/mL (19.17 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.9175 mL	9.5874 mL	19.1747 mL
	5 mM	0.3835 mL	1.9175 mL	3.8349 mL
	10 mM	0.1917 mL	0.9587 mL	1.9175 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: ≥ 4 mg/mL (7.67 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 4 mg/mL (7.67 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Thalidomide-NH-PEG4-COOH is an E3 ligase ligand-linker conjugate which can be used for synthesizing dCBP-1. dCBP-1 is a potent and selective heterobifunctional degrader of p300/CBP ^[1] .
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In Vitro	Thalidomide-NH-PEG4-COOH (S13) is the E3 ligase ligand-linker conjugate of dCBP-1 ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

1]. Raghu Vannam, et al. Targe	eted degradation of the enhancer ly	ysine acetyltransferases CBP and	d p300. Cell Chem Biol. 2020 Dec 31;S2451-	9456(20)30513-4.
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