

Product Data Sheet

Inhibitors

Screening Libraries

Proteins

Thalidomide-O-amido-PEG4-C2-NH2

 $\begin{array}{lll} \textbf{Cat. No.:} & HY\text{-}122710 \\ \\ \textbf{CAS No.:} & 1957236\text{-}22\text{-}4 \\ \\ \textbf{Molecular Formula:} & C_{25}H_{34}N_4O_{10} \\ \end{array}$

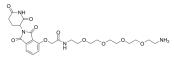
Molecular Weight: 550.56

Target: E3 Ligase Ligand-Linker Conjugates

Pathway: PROTAC

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.



BIOLOGICAL ACTIVITY

Description	Thalidomide-O-amido-PEG4-C2-NH2 is a synthesized E3 ligase ligand-linker conjugate that incorporates the Thalidomide based cereblon ligand and a linker used in PROTAC technology ^[1] .
IC ₅₀ & Target	Cereblon
In Vitro	PROTACs contain two different ligands connected by a linker; one is a ligand for an E3 ubiquitin ligase and the other is for the target protein. PROTACs exploit the intracellular ubiquitin-proteasome system to selectively degrade target proteins ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

 $[1]. \ James \ Bradner, et \ al. \ Methods \ to \ induce \ targeted \ protein \ degradation \ through \ bifunctional \ molecules. \ WO 2016 1055 18A1.$

[2]. Sato T, et al. Cereblon-Based Small-Molecule Compounds to Control Neural Stem Cell Proliferation in Regenerative Medicine. Front Cell Dev Biol. 2021;9:629326. Published 2021 Mar 11.

[3]. Nalawansha DA, et al. PROTACs: An Emerging Therapeutic Modality in Precision Medicine. Cell Chem Biol. 2020;27(8):998-985.

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