

## **Product** Data Sheet

Inhibitors

**Screening Libraries** 

Proteins

## Thalidomide-NH-PEG4-Ms

 $\begin{tabular}{lll} \textbf{Cat. No.:} & HY-131998 \\ \begin{tabular}{lll} \textbf{CAS No.:} & 2140807-24-3 \\ \begin{tabular}{lll} \textbf{Molecular Formula:} & $C_{22}H_{29}N_3O_{10}S$ \\ \end{tabular}$ 

Molecular Weight: 527.54

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-NH-PEG4-Ms is an E3 ligase ligand-linker conjugate that incorporates Thalidomide based cereblon ligand and a linker used for PROTAC BCL-XL degrader XZ739 <sup>[1]</sup> .
IC <sub>50</sub> & Target	Cereblon
In Vitro	XZ739, a CRBN-dependent PROTAC BCL-XL degrader with a DC <sub>50</sub> value of 2.5 nM in MOLT-4 cells after 16 h treatment. XZ739 also induces cell death through caspase-mediated apoptosis <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Xuan Zhang, et al. Discovery of PROTAC BCL-X L Degraders as Potent Anticancer Agents With Low On-Target Platelet Toxicity. Eur J Med Chem. 2020 Apr 15;192:112186.

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