

## **Product** Data Sheet

## (S)-Thalidomide-O-(1S,3r)-C4H4-N(Me)-Pip-C2-O-C-boc

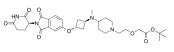
Cat. No.: HY-161183 Molecular Formula:  $C_{31}H_{42}N_4O_8$  Molecular Weight: 598.69

Target: Ligands for E3 Ligase; Autophagy

Pathway: PROTAC; Autophagy

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

Analysis.



## **BIOLOGICAL ACTIVITY**

Description	(S)-Thalidomide-O-(1S,3r)-C4H4-N(Me)-Pip-C2-O-C-boc is a conjugate of E3 ligase ligand and linker, consisting of Thalidomide (HY-14658) and the corresponding Linker. (S)-Thalidomide-O-(1S,3r)-C4H4-N(Me)-Pip-C2-O-C-boc can serve as a Cereblon ligand to recruit CRBN protein and serve as a key intermediate for the synthesis of complete PROTAC molecules.
IC <sub>50</sub> & Target	Cereblon (CRBN) <sup>[1]</sup>

## **REFERENCES**

[1]. Fischer ES, et al. Structure of the DDB1-CRBN E3 ubiquitin ligase in complex with thalidomide. Nature. 2014 Aug 7;512(7512):49-53.

[2]. Sun X, et al. Synergistic Inhibition of Thalidomide and Icotinib on Human Non-Small Cell Lung Carcinomas Through ERK and AKT Signaling. Med Sci Monit. 2018 May 15;24:3193-3203.

[3]. Bian C, et al. Thalidomide (THD) alleviates radiation induced lung fibrosis (RILF) via down-regulation of TGF-\(\beta\)/Smad3 signaling pathway in an Nrf2-dependent manner. Free Radic Biol Med. 2018 Dec;129:446-453.

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

Inhibitors

**Screening Libraries** 

**Proteins**