

## **Product** Data Sheet

Inhibitors

**Screening Libraries** 

## (1r,4r)-Thalidomide-piperidine-N(Me)-CH-cyclohexane-NH-Boc

Cat. No.: HY-163229 Molecular Formula:  $C_{31}H_{43}N_5O_6$  Molecular Weight: 581.7

Target: Ligands for E3 Ligase; Autophagy

Pathway: PROTAC; Autophagy

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

Analysis.

YOU NO NHOO

## **BIOLOGICAL ACTIVITY**

Description	(1r,4r)-Thalidomide-piperidine-N(Me)-CH-cyclohexane-NH-Boc is a conjugate of E3 ligase ligand and linker, consisting of Thalidomide (HY-14658) and the corresponding Linker. (1r,4r)-Thalidomide-piperidine-N(Me)-CH-cyclohexane-NH-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.

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