

Pomalidomide-amido-C1-Br

Chemical Properties

CAS No. : 2351106-38-0

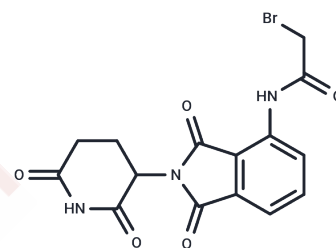
Formula: C₁₅H₁₂BrN₃O₅

Molecular Weight: 394.18

Keep away from direct sunlight

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.



Biological Description

Description	Pomalidomide-amido-C1-Br is a synthesized conjugate consisting of the Pomalidomide-based cereblon ligand and a linker, functioning as an E3 ligase ligand-linker. This compound serves as a tool for designing a B-Raf PROTAC degrader, specifically PROTAC B-Raf degrader 1, which exhibits anti-cancer activity[1].
Targets(IC50)	Others,E3 Ligase Ligand-Linker Conjugates,PROTAC Linker
In vitro	Pomalidomide-amido-C1-Br (Compound 6) serves as a precursor for synthesizing PROTAC B-Raf degrader 1 (Compound 2), an effective anticancer agent that induces apoptosis in cancer cells by accelerating B-Raf degradation via the ubiquitin-proteasome system, affecting Mcl-1 expression, a downstream protein associated with B-Raf, highlighting its mechanism of action[1].

Preparing Stock Solutions

	1mg	5mg	10mg
1 mM	2.5369 mL	12.6846 mL	25.3691 mL
5 mM	0.5074 mL	2.5369 mL	5.0738 mL
10 mM	0.2537 mL	1.2685 mL	2.5369 mL
50 mM	0.0507 mL	0.2537 mL	0.5074 mL

Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

Reference

Chen H, et al. Pomalidomide hybrids act as proteolysis targeting chimeras: Synthesis, anticancer activity and B-Raf degradation. Bioorg Chem. 2019 Jun;87:191-199.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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