

Catalog Number: CM19253

产品信息

Catalog Number:
CM19253

CAS号:
202591-23-9

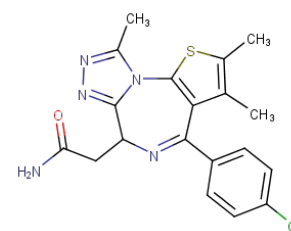
分子式:
C₁₉H₁₈ClN₅OS

主要靶点:
Epigenetic Reader Domain

主要通路:
表观遗传

分子量:
399.9

溶解度:
DMSO:200 mM



靶点活性

BET bromodomain:37 nM

体外活性

CPI203 inhibits BRD4 in vitro and in cells, while does not affect BRD4 kinase activity in vitro. [1] CPI203 exerts a cytostatic effect in all the 9 MCL cell lines analyzed with GI₅₀ ranging from 0.06 to 0.71 μM, with low cytotoxicity in normal PBMCs from healthy donors. Furthermore, lenalidomide and CPI203, by targeting IRF4 and MYC, efficiently activates the cell death program in MCL cells resistant to bortezomib. [2]

细胞实验

MCL primary cells and cell lines are incubated as indicated with lenalidomide and/or CPI203. MTT is added for 2-6 additional hours before spectrophotometric measurement. Each measurement is made in triplicate. Values are represented using untreated control cells. The GI₅₀ is calculated as the concentration that produced 50 % growth inhibition. Combination indexes (CIs) are calculated by using the Calcsyn software version 2.0. The interaction between two drugs is considered synergistic when CI < 1. (Only for Reference)

描述

CPI-203 is an effective BET bromodomain inhibitor (IC₅₀: 37 nM for BRD4).

储存

Powder: -20°C for 3 years | In solvent: -80°C for 2 years