

CPI203, BET bromodomain inhibitor

 Catalog
 Unit

 TBI4048-5MG
 5 mg

 TBI4048-25MG
 25 mg

Product Details

Formal Name: (S)-2-(4-(4-Chlorophenyl)-2,3,9-trimethyl-6h-Thieno[3,2-f][1,2,4]triazolo[4,3-a][1,4]diazepin-6-

yl)acetamide

Alternate Names: CPI-267203; 2-[(9S)-7-(4-Chlorophenyl)-4,5-13-trimethyl-3-thia-1,8,11,12-

tetrazatricyclo[8.3.0.02,6]trideca-2(6),4,7,10,12-pentaen-9-yl]acetamide

Molecular Formula: C₁₉H₁₈ClN₅OS

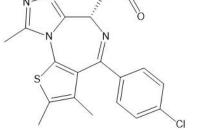
Formula Weight: 399.90 **CAS Number:** 1446144-04-2

Purity: >98%

Formulation: powder

Solubility: Soluble in DMSO (up to at least 25 mg/ml)

Storage: -20° C **Stability:** ≥ 1 year.



 NH_2

Applications

BET bromodomain inhibitor

Functions

CPI203, a JQ1 analog, is a potent and selective inhibitor of the BET bromodomain BRD4 (IC50 = 26 nM). It helps promote the maintenance of hematopoietic stem cells via repression of Myc expression. CPI203 reversibly suppressed intestinal stem cell differentiation in a mouse model. It enhanced the expansion of human cord blood hematopoietic stem cells without losing cell viability. CPI203 displays synergistic antitumor activity with various agents including lenalidomide, bortezomib, and PARP inhibitors.

Application Procedures

First dissolved in DMSO (up to at least 25 mg/ml), then diluted to aqueous buffer. Solutions in DMSO may be stored at -20° for up to 3 months.

For research use only.