
P5091

产品编号: D50869

CAS: 882257-11-6

分子式: C₁₂H₇Cl₂NO₃S₂

纯度: ≥98%

InChi: InChi=1S/C₁₂H₇Cl₂NO₃S₂/c1-6(16)10-5-8(15(17)18)12(20-10)19-9-4-2-3-7(13)11(9)14/h2-5H, 1H3

InChi Key: LKZLGMAAKNEGCH-UHFFFAOYSA-N

Smiles: CC(=O)C1=CC(=C(SC2C=CC=C(Cl)C=2Cl)S1)[N+]([O-])=O

外观: 固体粉末

作用通路: Deubiquitinase

溶解性: DMSO up to 100 mM

保存条件: Store in dry, dark place for one year.

产品介绍: P5091 (P005091) is a novel potent and selective inhibitor of ubiquitin-specific protease 7 (USP7) with an IC₅₀ ~4.2 μM. It does not inhibit other DUBs or other families of cysteine proteases tested (EC₅₀ > 100 mM). It induces apoptosis in MM cells resistant to conventional and bortezomib therapies. Biochemical and genetic studies showed that blockade of HDM2 and p21 abrogated P5091-induced cytotoxicity. In animal tumor model studies, P5091 was well tolerated, and it inhibited tumor growth, and prolongs survival. Combining P5091 with lenalidomide, HDAC inhibitor SAHA, or dexamethasone triggered synergistic anti-MM activity.