For Research Use Only ML-SA1



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Catalog Number: CM06793

产品信息

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CM06793 CAS号:

332382-54-4

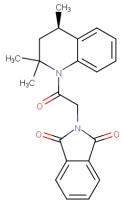
分子式: C₂₂H₂₂N₂O₃

主要靶点:

Others | TRP/TRPV Channel

主要通路: 离子通道|其他 分子量: 362.42 溶解度:

DMSO:17 mg/mL (46.90 mM)



靶点活性

ZIKV:52.99 μ M (IC50)|DENV2:8.3 μ M (IC50)

体外活性

ML-SA1 (25 μ M; 0~14 hours; A549 cells) possibly affects the entry of DENV2 into host cells[1]. ML-SA1 (0~200 μ M; A549 cells) shows that there is no cytotoxicity to the cell line observed, even at concentrations up to 200 μ M. ML-SA1 (0~50 μ M; A549 cells) significantly suppresses DENV2 at the RNA levels and the IC50 is 8.93 μ M[1]. ML-SA1 results in a dose-dependent decrease in ZIKV in A549 cells at both the RNA and protein levels, and the IC50 value of ML-SA1 against ZIKV RNA is 52.99 μ M. ML-SA1, as an activator of TRPMLs, appears to be a potent inhibitor of DENV2 and ZIKV in vitro. ML-SA1 promotes lysosomal acidification and protease activity to inhibit viral infection. ML-SA1 can induce autophagy in Huh7 cells or A549 cells[1].

描述

ML-SA1 (Mucolipin synthetic agonist 1) is a selective TRPML agonist, inhibits Dengue virus 2 (DENV2) and Zika virus (ZIKV) by promoting Lysosomal acidification and protease activity. The IC50 value of ML-SA1 against DENV2 RNA and ZIKV RNA is 8.3 μ M and 52.99 μ M, respectively[1].

储存

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