For Research Use Only **Devimistat**



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Catalog Number: CM03649

产品信息

Catalog Number: CM03649 分子量: 388.59 CAS号: 溶解度:

95809-78-2 DMSO:78 mg/mL (200.72 mM),Ethanol:78 mg/mL (200.72 mM),H2O:<1 mg/mL 分子式: C₂₂H₂₈O₂S₂

主要靶点: Apoptosis|Dehydrogenase|Mitochondrial Metabolism

主要通路: 代谢|凋亡

体外活性

In vitro, CPI-613 produces the selective toxicity against several tumor cell lines including H460 human lung cancer cells and Saos-2 human sarcoma cells with EC50 of 120 $\,\mu$ M and 120 $\,\mu$ M, respectively. CPI-613 disrupts H460 cancer cell mitochondrial metabolism including inhibition of PDH complex activity and loss of mitochondrial membrane potential in a time- and drug dose-dependent fashion. In addition, CPI-613 (240 $\,\mu$ M) also induces both apoptotic and non-apoptotic cell death in H460 human lung cancer and Saos-2 human sarcoma cells. [1]

体内活性

CPI-613 (25 mg/kg) has potent anticancer activity in a human tumor xenograft model of of a pancreatic tumor cell (BxPC-3). Similarly, CPI-613 (10 mg/kg) also produces significant tumor growth inhibition of H460 human non-small cell lung carcinoma in mouse model. Besides, CPI-613 produces little or no side-effect toxicity in expected therapeutic dose ranges in large animal models and has the maximum tolerated dose of 100 mg/kg in mice. [1]

CPI-613, a lipoate analog, inhibits mitochondrial enzymes pyruvate dehydrogenase (PDH) and $\,^{\alpha}$ -ketoglutarate dehydrogenase, disrupts tumor cell mitochondrial metabolism. It has potential chemopreventive and antineoplastic activities, and has been used in trials studying the treatment of Cancer, Lymphoma, Solid Tumors, Advanced Cancer, and Pancreatic Cancer, among others.

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