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Catalog Number: CM04468

产品信息

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CAS号: 927822-86-4

分子式: C₁₆H₁₆Cl₄N₂O₄S₄

HIF|HIF/HIF Prolyl-Hydroxylase

主要通路: 表观遗传|血管生成|代谢

靶点活性

HIF-1 α :20 μ M

KC7F2 inhibits HRE-driven transcription and decreases HIF-1 lpha protein levels in LN229-HRE-AP cells. KC7F2 shows a dose-response cytotoxicity with IC50 of approximately 15 to 25 $^{\mu}$ M in cancer cells MCF7, LNZ308, A549, U251 mg, and LN229. In D54 mg glioma cells, KC7F2 inhibits colony formation, especially under hypoxia. [1] In hypoxic microglial cultures, KC7F2 downregulates the expression of TfR and DMT, and reduces the HIF-1 $^{\alpha}$ mediated iron accumulation. [2]

DMSO:57 mg/mL (100 mM)

KC7F2 significantly reduces the latent period in the pentylenetetrazole kindling rat model and increases the rate of spontaneous recurrent seizures during the chronic stage in the lithium-pilocarpine rat model. [3]

分子量: 570.38

溶解度:

细胞实验

Cells are seeded onto 96-well plates (4 × 103/well) and cultured under normoxic (21% O2) and hypoxic (1% O2) conditions with different concentrations of KC7F2 for 72 h or treated for various times with 20 μ M KC7F2. For proliferation analysis, cells are fixed with 50% trichloroacetic acid for 1 h at 4°C, followed by staining with 0.4% sulforhodamine B dissolved in 1% acetic acid for 30 min at room temperature. Plates are washed five times with 1% acetic acid to remove unbound dye. Bound dye is dissolved by adding 10 mM unbuffered Tris base. Cell proliferation is calculated by measuring OD values at 564 nm using a spectrophotometer.(Only for Reference)

描述

KC7F2 is a potent HIF-1 pathway inhibitor with potential anti-cancer activity.

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