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Catalog Number: CM04540

产品信息

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CAS号: 1423715-09-6

分子式: C₁₉H₂₀ClN₃O₅S

主要靶点:

Histone Demethylase|Apoptosis

主要通路: 表观遗传|凋亡

分子量: 437.9 溶解度:

H2O:<1 mg/mL,DMSO:36 mg/mL (82.2 mM),Ethanol:<1 mg/mL

靶点活性

LSD1:13 nM

体外活性

在负荷OCI-AML3异种移植物的小鼠中,SP2509(25 mg/kg i.p.),能够抑制肿瘤生长,延迟动物的存活.

体内活性

在OCI-AML3中,SP250能够抑制集落形成,诱导细胞凋亡。在AML细胞中,SP2509能够抑制LSD1与CoREST作用,增加启动子特异性 H3K4Me3,同时诱导p53,p21和C/EBP α 。在原代AML细胞中,SP2509诱导细胞的增殖。

细胞实验

Cultured AML cells are treated with SP2509 and/or PS for 96?h. At the end of treatment, cells are washed free of the drugs and 500 cells per condition are plated in methylcellulose and incubated at 37?°C. Colony formation is measured 7–10 days after plating.(Only for Reference)

描述

SP2509 is a specific histone demethylase LSD1 inhibitor(IC50 =13 nM).

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