

Catalog Number: CM03745

产品信息

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CM03745

CAS号:
14892-97-8

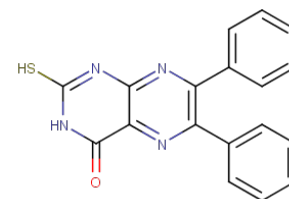
分子式:
C₁₈H₁₂N₄OS

主要靶点:
Apoptosis|CRISPR/Cas9|DNA/RNA
Synthesis

主要通路:
DNA 损伤和修复|细胞周期|DNA 损伤
和修复|凋亡

分子量:
332.38

溶解度:
DMSO:66 mg/mL (198.57
mM);H₂O:< 1 mg/mL (insoluble
or slightly soluble);Ethanol:3
mg/mL (9.03 mM)



体外活性

SCR7在200 μM及以上浓度下有效抑制多聚体的形成。SCR7成功抑制了MCF7、A549、HeLa、T47D、A2780、HT1080和Nalm6细胞的增殖，其IC₅₀分别为40、34、44、8.5、120、10和50 μM。[1] SCR7抑制了CRISPR-Cas9诱导的DSBs的NHEJ修复。[2]

体内活性

SCR7处理（10 mg/kg，肌肉注射）显著减少了乳腺癌诱导的肿瘤，并且与对照组相比，生命期增加了4倍。然而，在携带Dalton's淋巴瘤肿瘤模型的瑞士白鼠中，SCR7（20 mg/kg，腹腔注射）既不表现出肿瘤退缩也不增加生命期。在BALB/c小鼠中，SCR7（20 mg/kg，腹腔注射）显著增强了辐射、依托泊苷和3-氨基苯甲酰胺对源自Dalton's淋巴瘤（DLA）细胞的肿瘤的细胞毒作用。[1]

细胞实验

Cell proliferation of cancer cells are determined by MTT and trypan blue assays. Briefly, MCF7, CEM, HeLa, A549, HT1080, A2780, T47D, Nalm6, N114 and K562 cells are grown in presence of SCR7 (10, 50, 100, and 250 μM) for 24 or 48 h, and subjected to MTT or trypan blue assays. Each experiment is repeated a minimum of three independent times. (Only for Reference)

储存

Powder: -20°C for 3 years | In solvent: -80°C for 1 year | Shipping with blue ice.