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Catalog Number: CM06124

产品信息

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CAS号: 880635-03-0

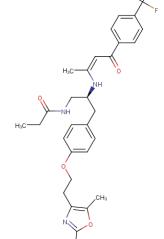
分子式: C₃₅H₃₆F₃N₃O₄

主要靶点: PPAR

主要通路: DNA损伤和修复|代谢

分子量: 619.67 溶解度:

DMSO:120 mg/mL (193.65 mM),Sonification is recommended.



体外活性

A specific PPAR α antagonist, GW6471, induced both apoptosis and cell cycle arrest at GO/G1 in VHL(+) and VHL(-) RCC cell lines (786-O and Caki-1) associated with attenuation of the cell cycle regulatory proteins c-Myc, Cyclin D1, and CDK4;?this data was confirmed as specific to PPAR α antagonism by siRNA methods.

GW6471 is an antagonist of PPAR α with IC50 of 0.24 μ M. GW6471 enhances the binding affinity of the PPAR α ligand-binding domain to the co-repressor proteins SMRT and NCoR.

储存

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