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## Entinostat (MS-275)

产品编号: D50840

CAS: 209783-80-2

分子式: C<sub>21</sub>H<sub>20</sub>N<sub>4</sub>O<sub>3</sub>

纯度: ≥98%

InChi: InChI=1S/C<sub>21</sub>H<sub>20</sub>N<sub>4</sub>O<sub>3</sub>/c22-18-5-1-2-6-19(18)25-20(26)17-9-7-15(8-10-17)13-24-21(27)28-14-16-4-3-11-23-12-16/h1-12H,13-14,22H<sub>2</sub>, (H,24,27)(H,25,26)

InChi Key: INVTYAOGFAGBOE-UHFFFAOYSA-N

Smiles: NC1C=CC=CC=1NC(=O)C1C=CC(CNC(=O)OCC2=CC=CN=C2)=CC=1

外观: 固体粉末

作用通路: Apoptosis

溶解性: DMSO up to 100 mM

保存条件: Store in dry, dark place for one year.

产品介绍: Entinostat (MS-275) is a potent and selective HDAC inhibitor. It inhibits HDAC1 and HDAC3 with IC<sub>50</sub> of 0.51 μM and 1.7 μM, but not the other HDACs 4, 6, 8, and 10 (IC<sub>50</sub> > 100 μM). MS-275 induces accumulation of p21WAF1/CIP1 and gelsolin in K562 cell. It was shown to reduce S-phase cells and induce G1-phase cells in A2780 cells. MS-275 shows great inhibition to human leukemia and lymphoma cells, decreases expression of cyclin D1 and the anti-apoptotic proteins Mcl-1 and XIAP. In vivo MS-275 exhibits great antitumor activity against human tumor xenografts. Currently it is in Phase II/III clinical trials for Hodgkin's lymphoma and advanced breast cancer.