

FK-228(Romidepsin)

Cat. No. CEI-0370

Lot. No. (See product label)

Introduction

Description

Unlike TSA, the active form redFK of Romidepsin strongly inhibits HDAC1 and HDAC2 with IC₅₀ of 1.6 nM and 3.9 nM, respectively, but is relatively weak in inhibiting HDAC4 and HDAC6 with IC₅₀ 25 nM and 790 nM, respectively. Romidepsin is 17-23 times weaker than redFK in inhibiting these HDACs with IC₅₀ of 36 nM, 47 nM, 510 nM, and 14 μ M, respectively.

Product Information

CAS No.	128517-07-7
Molecular Formula	C ₂₄ H ₃₆ N ₄ O ₆ S ₂
Chemical Name	Cyclo[(2Z)-2-amino-2-butenoyl-L-valyl-(3S,4E)-3-hydroxy-7-mercapto-4-heptenoyl-D-valyl-D-cysteinyl], cyclic (3→5)-disulfide
Molecular Weight	540.7
Purity	>99%
Targets	HDAC
Solubility	DMSO 10 mg/mL, Water

Storage and Shipping Information

Storage	2 years at -20 centigrade
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