

Catalog # 10-2365 Romidepsin

128517-07-7 FK228, FR901228, Depsipeptide Lot # X106426

A potent and selective inhibitor of class I histone deacetylases (HDACs), IC₅₀ = 36, 47, 510 and 14,000 nM for HDAC1, 2, 4 and HDAC6, respectively.¹ Induces apoptosis in a variety of cell lines² and displays antitumor activity in mouse models³. Recently approved for clinical use in T-cell lymphoma.⁴

- 1) Furumai et al. (2002), FK228 (depsipeptide) as a natural prodrug that inhibits class I histone deacetylases; Cancer Res.., **62** 4916
- Panicker et al. (2010), Romidepsin (FK228/depsipeptide) controls growth and induces apoptosis in neuroblastoma tumor cells; Cell Cycle, 9 1830
- 3) Ueda et al. (1994), FR901228, a novel anti-tumor bicyclic depsipeptide produced by Chromobacterium violaceum No. 968. III. Antitumor activities on experimental tumors in mice; J. Antibiot. (Tokyo), **47** 315
- 4) VanderMolin et al. (2011), Romidepsin (Istodax, NSC 630176, FR901228, FK228, depsipeptide): a natural product recently approved for cutaneous T-cell lymphoma; J. Antibiotic. (Tokyo), **64** 525

PHYSICAL DATA

Molecular Weight: 540.71

Molecular Formula: C₂₄H₃₆N₄O₆S₂ Purity: 98% by HPLC

NMR: (Conforms)

Solubility: DMSO (up to 10 mg/ml)

Physical Description: Colorless waxy solid – vials may appear empty

Storage and Stability: Store as supplied desiccated at -20°C for up to 2 years from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 1 month.

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