

Salermide



Orders: 877-616-CELL (2355)

orders@cellsignal.com

Support: 877-678-TECH (8324)

Web: info@cellsignal.com

cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

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For Research Use Only. Not for Use in Diagnostic Procedures.

Background

The reverse amide Salermide is a strong inhibitor of the sirtuin proteins SirT1 and SirT2. Sirtuins are nicotinamide adenine dinucleotide-dependent protein deacetylases involved in cell aging and lifespan regulation. Nuclear SirT1 is implicated in regulating apoptosis, cellular senescence, aging, and longevity; SirT2 is involved in cytoskeletal regulation and progression through mitosis. Exposure of human cancer cell lines to Salermide results in the reactivation of proapoptotic genes repressed by SirT1 and tumor-specific cell death (1). Salermide treatment of human breast cancer cells leads to decreased SirT1 expression and increased acetylation and activation of p53 (2). Inhibition of both SirT1 and SirT2 by Salermide is required to induce p53 acetylation and cell death (3). Salermide treatment of human pancreatic cancer cells potentiates the anticancer effects of a cytotoxin, reducing pancreatic cancer cell progression and stopping the cell cycle at G1 (4).

Molecular Formula $C_{26}H_{22}N_2O_2$ **Molecular Weight** 394.5 g/mol

Purity >98%

CAS 1105698-15-4

Solubility Soluble in DMSO at 40 mg/mL, ethanol at 10 mg/mL, or DMF at 30 mg/mL.

Store lyophilized at -20°C, desiccated. In lyophilized form, the chemical is stable for 24 months. Once in Storage

solution, store at -20°C and use within 1 month to prevent loss of potency. Aliquot to avoid multiple

freeze/thaw cycles.

Directions for Use Salermide is supplied as a lyophilized powder. For a 15 mM stock, reconstitute 5 mg of powder in 0.84

mL of DMSO. Working concentrations and length of treatment can vary depending on the desired

effect.

Background References 1. Lara, E. et al. (2009) Oncogene 28, 781-91.

2. Dastjerdi, M.N. et al. (2013) Res Pharm Sci 8, 79-89.

3. Peck, B. et al. (2010) Mol Cancer Ther 9, 844-55.

4. Yar Saglam, A.S. et al. (2016) EXCLI J 15, 246-55.

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