

Tubulin/HDAC-IN-4

Chemical Properties

CAS No. :

Formula: C₂₄H₂₆N₂O₆

Molecular Weight:

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Actual storage temperature shall be subject to the COA.

Biological Description

Description	Tubulin/HDAC-IN-4 (compound 9n) is a dual inhibitor targeting microtubules (Tubulin) and HDAC, exhibiting IC ₅₀ values of 0.73, 0.43, 0.62, and 2.34 μ M for HDAC1, HDAC2, HDAC6, and HDAC7, respectively. It disrupts microtubule assembly by binding to the colchicine site and induces apoptosis and G2/M cell cycle arrest. Additionally, Tubulin/HDAC-IN-4 significantly increases intracellular ROS levels and demonstrates both anti-angiogenic and anticancer properties.
Targets(IC50)	Apoptosis, Reactive Oxygen Species, Microtubule Associated, HDAC
In vitro	Tubulin/HDAC-IN-4 (compound 9n), in concentrations ranging from 0 to 10 μ M over 72 hours, exhibits cytotoxic effects against MDA-MB-231 and A549 cells, with IC ₅₀ values of 0.34, 0.29, 0.016, 0.15, and 0.16 μ M for PC-3, U251, and MCF-7 cells, respectively. The compound also inhibits colony formation in PC-3 cells in a dose-dependent manner at concentrations of 2.5, 5, 10, 20, and 40 nM over 24 hours. Furthermore, Tubulin/HDAC-IN-4 disrupts tubulin polymerization at concentrations of 0.2, 1, 5, and 25 μ M with an IC ₅₀ of 4.82 μ M. It increases the expression of Ac- α -tubulin and Ac-Histone H3 in PC-3 cells at 0.08, 0.16, and 0.32 μ M over 24 hours. Additionally, the compound induces apoptosis, G2/M phase cell cycle arrest, and significantly elevates intracellular ROS levels at 0.08, 0.16, and 0.32 μ M over the same period.
In vivo	Administered intravenously at doses of 10 and 20 mg/kg every two days for 21 days, Tubulin/HDAC-IN-4 demonstrates anticancer activity.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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