

Product Name : HDAC-IN-57

Synonyms

Cat No. : M36851

CAS Number : 2716217-79-5

Molecular Formula : C21H19N3O4

Formula Weight : 377.39

Chemical Name

HDAC-IN-57 is an orally active inhibitor of histone deacetylases (HDAC), with IC50s of 2.07 nM, 4.71 nM, 2.4 nM and 107 \cdot nM for HDAC1, HDAC2, HDAC6, HDAC8, respectively. HDAC-IN-57 can inhibits LSD1, with IC50 of 1.34 μ M.HDAC-IN-57 Description

induces apoptosis, and has anti-tumor activity.

Pathway : Cell Cycle/DNA Damage

Target : HDAC

Receptor : HDAC | Apoptosis

Solubility

SMILES $: \quad \mathsf{C}(\mathsf{NCC1} = \mathsf{CC} = \mathsf{C}(\mathsf{C}(\mathsf{NO}) = \mathsf{O})\mathsf{C} = \mathsf{C1})(=\mathsf{O})\mathsf{C} = 2\mathsf{C} = \mathsf{C}(\mathsf{N} = \mathsf{CC2})\mathsf{C3} = \mathsf{CC} = \mathsf{C}(\mathsf{OC})\mathsf{C} = \mathsf{C3}$

: (-20℃) Storage

Stability : ≥2 years

Reference

^{1.} Duan Y, et al. Discovery of novel, potent, and orally bioavailable HDACs inhibitors with LSDI inhibitory activity for the treatment of solid tumors. Eur J Med Chem. 2023 Jun 5;254:115367.?