

STING-IN-13

Chemical Properties

CAS No. :

Formula:

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	STING-IN-13 is a selective STING inhibitor that effectively suppresses downstream signaling of the STING pathway and STING-mediated inflammation. It exhibits low toxicity and is suitable for research related to STING-associated inflammatory and autoimmune diseases.
Targets(IC50)	Interleukin,STING
In vitro	STING-IN-13 (Compound HY2) at a concentration of 30 μ M interacts directly with STING in RAW264.7 cells. Pre-treatment with STING-IN-13 (0.03 μ M-0.3 μ M for 1 hour, followed by co-incubation for 2-4 hours) effectively inhibits downstream signaling in the STING pathway induced by SR717 in THP1 and RAW264.7 cells, significantly reducing the production of inflammatory factors IFN- β and CXCL10. At concentrations of 0.4 μ M-3.3 μ M over 48 hours, STING-IN-13 demonstrates excellent safety with low cytotoxicity in THP1 cells. Additionally, STING-IN-13 (5 μ M, pre-treatment for 1 hour followed by treatment for 2 hours) specifically inhibits STING-driven IFN β expression in THP1 cells without affecting the TLR pathway.
In vivo	STING-IN-13 (Compound HY2) (10,20 mg/kg, i.p. one dose) enhances survival rates and protects renal and hepatic function in a cisplatin-induced AKI mouse model, while also reducing tubular damage and inflammatory cell infiltration.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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