

QNX-10

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	QNX-10 is a fatty acid synthase (FASN) inhibitor with anticancer properties (IC ₅₀ = 0.7 μM). It exhibits potent FASN inhibition and cytotoxicity against colorectal and breast cancer cells. By upregulating the pro-apoptotic protein Bax and downregulating the anti-apoptotic protein Bcl-xL, QNX-10 induces apoptosis and causes cell cycle arrest at the S phase. This compound is utilized for investigating anticancer therapies targeting the FASN enzyme.
Targets(IC ₅₀)	Apoptosis,Bcl-2 Family,Fatty Acid Synthase
In vitro	QNX-10, administered at concentrations ranging from 0-10 μM for 48 hours, exhibits cytotoxicity against HCT-116, Caco-2, and MCF-7 cells, with IC ₅₀ values of 1.5 μM, 2.1 μM, and 1.8 μM, respectively. It is non-toxic to HEK-293 cells, with an IC ₅₀ of 30.4 μM. QNX-10 (0.5-2.5 μM, 48 hours) inhibits cell cycle progression by inducing S-phase cell cycle arrest in HCT-116 cells. Additionally, QNX-10 (0.5-2.5 μM) shows dose-dependent apoptotic characteristics by increasing Bax and decreasing Bcl-xL and FASN levels in HCT-116 cells.

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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