
Product Data Sheet

Product Name: PI3K/Akt/mTOR-IN-2

Cat. No.: GC66463

Chemical Properties

Cas. No. 2757804-89-8

Formula $C_{17}H_{13}F_2NO$ M.Wt 285.29

Solubility DMSO : ≥ 100 mg/mL (350.52 mM) Storage Store at $-20^{\circ}C$

General tips For obtaining a higher solubility , please warm the tube at $37^{\circ}C$ and shake it in the ultrasonic bath for a while. Stock solution can be stored below $-20^{\circ}C$ for several months.

Shipping Condition Evaluation sample solution : ship with blue ice All other available size: ship with RT , or blue ice upon request.

Structure

Background

PI3K/Akt/mTOR-IN-2 is a **PI3K/AKT/mTOR** pathway inhibitor. PI3K/Akt/mTOR-IN-2 possess anti-cancer effects and selectivity against MDA-MB-231 cells with **IC₅₀** value of $2.29 \mu M$. PI3K/Akt/mTOR-IN-2 can induce cancer cell cycle arrest and **apoptosis**^[1].

PI3K/Akt/mTOR-IN-2 (compound 23) ($0.5 - 100 \mu M$; 72 hours) exhibits effective anti-cancer activity with **IC₅₀**s of $2.29 - 24.63 \mu M$, of which, **IC₅₀** in MDA-MB-231 is $2.29 \mu M$ ^[1].

PI3K/Akt/mTOR-IN-2 ($1 \mu M$, $2 \mu M$ and $4 \mu M$; 24 hours) induces growth inhibition of MDA-MB-231 cells by cell cycle arrest at G₀/G₁^[1].

PI3K/Akt/mTOR-IN-2 ($1 \mu M$, $2 \mu M$ and $4 \mu M$; 24, 48 and 72 hours) induces apoptosis in MDA-MB-231 cells with both dose- and time-dependent manners^[1].

PI3K/Akt/mTOR-IN-2 ($1 \mu M$, $2 \mu M$ and $4 \mu M$; 48 hours) increases the expression of Bax, and decreases the expression of Bcl-2 in MDA-MB-231 cells^[1].

PI3K/Akt/mTOR-IN-2 ($1 \mu M$, $2 \mu M$ and $4 \mu M$; 24 hours) induces mitochondria-dependent apoptosis in MDA-MB-231 cells through disruption of MMP, accumulation of ROS, depletion of GSH and elevation of intracellular Ca^{2+} ^[1].

Caution: Product has not been fully validated for medical applications. For research use only.

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Cell Proliferation Assay

Cell Line: PC3, BGC-823, A549, MCF-7, MDA-MB-231 cells and MCF-10A cells^[1]

Concentration: 0.5 - 100 μ M

Incubation
Time: 72 hours

Result: Exhibited effective anti-cancer activity with IC₅₀s of 2.29 - 24.63 μ M, of which, IC₅₀ in MDA-MB-231 was 2.29 μ M.

Cell Cycle Analysis

Cell Line: MDA-MB-231^[1]

Concentration: 1 μ M, 2 μ M and 4 μ M

Incubation
Time: 24 hours

Result: Induced growth inhibition of MDA-MB-231 cells by cell cycle arrest at G₀/G₁.

Apoptosis Analysis

Cell Line: MDA-MB-231^[1]

Concentration: 1 μ M, 2 μ M and 4 μ M

Incubation
Time: 24, 48 and 72 hours

Result: Induced apoptosis in MDA-MB-231 cells with both dose- and time-dependent manners.

Western Blot Analysis

Cell Line: MDA-MB-231^[1]

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Concentration: 1 μ M, 2 μ M and 4 μ M

Incubation
Time: 48 hours

Result: Increased the expression of Bax, and decreased the expression of Bcl-2

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