Product Data Sheet

Isochamaejasmin

Cat. No.: HY-N3497 CAS No.: 93859-63-3 Molecular Formula: $C_{30}H_{22}O_{10}$ 542.49 Molecular Weight:

Target: DNA/RNA Synthesis; NF-κB; Parasite; Apoptosis

Pathway: Cell Cycle/DNA Damage; NF-кВ; Anti-infection; Apoptosis

Please store the product under the recommended conditions in the Certificate of Storage:

Analysis.

BIOLOGICAL ACTIVITY

Description

Isochamaejasmin is a biflavonoid with anti-cancer, antiplasmodial and insecticidal activities. Isochamaejasmin displays a potent NF-κB (NF-κB) activation activity. Isochamaejasmin could cause DNA damage and induce apoptosis via the mitochondrial pathway in AW1 cells $^{[1][2]}$. Isochamaejasmin also has a moderate antiplasmodial activity (IC₅₀ of 7.3 μ M for P. falciparum) and relatively low cytotoxicity (CC_{50} of 29.0 μ M)^[3].

In Vitro

Isochamaejasmin (6.25-100 µM; 24-72 h) shows potential toxicity against AW1 cells via time- and dose-dependent manners. Isochamaejasmin (1000 mg/L, 500 mg/L, 250 mg/L, 125 mg/L, 62.5 mg/L; 24 h, 48 h, 72 h, and 96 h) has potential toxicity against H. zea larvae via time- and dose-dependent manners^[1].

Isochamaejasmin (40-80 µM; 24 h) causes DNA damage and increases the levels of yH2AX and OGG1 in AW1 cells. The cell cycle is arrested at the G2/M phase^[1].

Isochamaejasmin (20-80 μ M; 24 h) induces apoptosis of AW1 cells in a dose-dependent manner [1].

Isochamaejasmin (20-80 μM; 24 h) shows decline in the MMP, upregulation of Bax/Bcl-2 expression resulting in the release of cytochrome c into the cytosol, activation of caspase-3/9, and cleavage of PARP^[1].

Isochamaejasmin shows a dose-dependent rise in the reactive oxygen species (ROS) levels, accumulation of a lipid peroxidation product, and inactivation of antioxidant enzymes in AW1 cells^[1].

Isochamaejasmin induces the expression of a NF- κ B-directed reporter gene in transfected HeLa cells with an EC₅₀ of 3.23 μ M. The Isochamaejasmin-stimulated NF-кВ reporter activity is accompanied by nuclear translocation of NF-кВ proteins and is blocked by a dominant-negative construct of IκBα. Isochamaejasmin also induces time-dependent phosphorylation of the mitogen-activated protein kinases ERK1/2 and p38, and PKC $\delta^{[2]}$.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Viability Assay^[1]

Cell Line:	Larvae and neuronal cells (AW1)
Concentration:	6.25 μM, 12.5 μM, 25 μM, 50 μM and 100 μM
Incubation Time:	24 h, 48 h and 72 h
Result:	Had potential toxicity against H. zea both in vivo and in vitro via time- and dose-dependent manners.
Cell Cycle Analysis ^[1]	
Cell Line:	Neuronal cells (AW1)

Concentration:	40 μM and 80 μM
Incubation Time:	24 h
Result:	The cell cycle was arrested at the G2/M phase.
Apoptosis Analysis ^[1]	
Cell Line:	Neuronal cells (AW1)
Concentration:	20 μM, 40 μM, and 80 μM
Incubation Time:	24 h
Result:	Induced apoptosis via the mitochondrial pathway in AW1 cells.
Western Blot Analysis ^[1]	
Cell Line:	Neuronal cells (AW1)
Concentration:	20 μM, 40 μM, and 80 μM
Incubation Time:	24 h
Result:	Showed upregulation of Bax/Bcl-2 expression resulting in the release of cytochrome c into the cytosol, activation of caspase-3/9, and cleavage of PARP.

REFERENCES

- [1]. Yuanhang Ren, et al. Isochamaejasmin induces toxic effects on Helicoverpa zea via DNA damage and mitochondria-associated apoptosis. Pest Manag Sci. 2021 Jan;77(1):557-567.
- [2]. Qinghai Tian, et al. Stereospecific induction of nuclear factor-kappaB activation by isochamaejasmin. Mol Pharmacol. 2005 Dec;68(6):1534-42.
- [3]. Liene Dhooghe, et al. Antiplasmodial activity of (I-3,II-3)-biflavonoids and other constituents from Ormocarpum kirkii. Phytochemistry. 2010 May;71(7):785-91.

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

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