Proteins

Screening Libraries

Product Data Sheet

Niacin hydrochloride

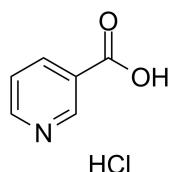
Cat. No.: HY-B0143A CAS No.: 636-79-3 Molecular Formula: C₆H₆ClNO₂

Molecular Weight: 159.57

Target: Endogenous Metabolite; Apoptosis Pathway: Metabolic Enzyme/Protease; Apoptosis

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

Analysis.



BIOLOGICAL ACTIVITY

Description

Niacin (Vitamin B3; Nicotinic acid) hydrochloride is an orally active B3 vitamin that is an essential nutrient for humans. Niacin hydrochloride plays a key role in energy metabolism, cell signaling cascades regulating gene expression and apoptosis. Niacin hydrochloride is also used in the study of cardiovascular diseases^{[1][2]}.

IC₅₀ & Target

Microbial Metabolite

Human Endogenous Metabolite

In Vitro

Niacin hydrochloride (0-900 µM, 42 hours) significantly increases GSH levels and decreases ROS levels, and affects the expression of genes related to apoptosis and lipid metabolism^[1].

Niacin hydrochloride (0-40 µM, 24 hours) can inhibit cancer invasive activity at low dose but with no influence on

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

RT-PCR^[1]

Cell Line:	Cumulus cells and oocytes of prepubertal gilts
Concentration:	600 μΜ
Incubation Time:	42 hours
Result:	Up-regulated the relative expression of the anti-apoptotic gene BCL2 and lipid metabolism gene ACACA while down-regulated the pro-apoptotic gene BAX.

Cell Proliferation Assay^[2]

Cell Line:	Rat ascites hepatoma cell line of AH109A
Concentration:	0-40 μΜ
Incubation Time:	24 hours
Result:	Had no effect on the proliferation of AH109A cells but suppressed cell invasion from 2.5 μM to 40 $\mu\text{M}.$

In Vivo

Niacin hydrochloride (subcutaneous injection, 3-300 mg/kg, once) can induce vasodilation in a dose-dependent manner within minutes in male C57BL/6 mice^[3].

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Animal Model:	Male C57BL/6 mice ^[3]
Dosage:	3-300 mg/kg
Administration:	Subcutaneous injection; once
Result:	Induced vasodilation in a dose-dependent manner.

CUSTOMER VALIDATION

- Mil Med Res. 2022 Aug 23;9(1):46.
- Mol Cell. 2023 Aug 11;S1097-2765(23)00605-6.
- Gut Microbes. 2023 Jan-Dec;15(1):2186114.
- J Nanobiotechnology. 2022 Mar 9;20(1):120.
- Glia. 2018 Feb;66(2):256-278.

See more customer validations on www.MedChemExpress.com

REFERENCES

- [1]. Areeg M Almubarak, et al. Supplementation with Niacin during in vitro maturation improves the quality of porcine embryos. Theriogenology. 2021 Jul 15;169:36-46. doi: 10.1016/j.theriogenology.2021.04.005. Epub 2021 Apr 18.
- [2]. Nobuhiro Hirakawa, et al. Anti-invasive activity of niacin and trigonelline against cancer cells. Biosci Biotechnol Biochem. 2005 Mar;69(3):653-8.
- [3]. Kang Cheng, et al. Antagonism of the prostaglandin D2 receptor 1 suppresses nicotinic acid-induced vasodilation in mice and humans. Proc Natl Acad Sci U S A. 2006 Apr 25;103(17):6682-7.

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

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