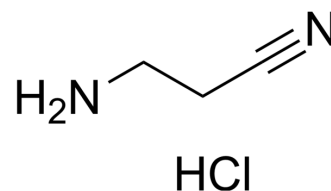


β-Aminopropionitrile hydrochloride

Cat. No.:	HY-Y1750A
CAS No.:	646-03-7
Molecular Formula:	C ₃ H ₇ ClN ₂
Molecular Weight:	107
Target:	Monoamine Oxidase; Endogenous Metabolite
Pathway:	Neuronal Signaling; Metabolic Enzyme/Protease
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	β-Aminopropionitrile (BAPN) hydrochloride is a specific, irreversible and orally active lysyl oxidase (LOX) inhibitor. β-Aminopropionitrile hydrochloride targets the active site of LOX or LOXL isoenzymes ^{[1][2]} .																		
IC₅₀ & Target	Lysyl Oxidase																		
In Vitro	<p>β-Aminopropionitrile (BAPN) normalizes the expression of GLUT4 and adiponectin, and improves glucose uptake in an in vitro model of insulin resistance^[1].</p> <p>β-Aminopropionitrile (500 μM; 72 h) blocks the hypoxia-induced EMT morphological and marker protein changes, and inhibits invasion and migration capacities of cervical carcinoma cells in vitro^[2].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>3T3-L1 adipocytes</td> </tr> <tr> <td>Concentration:</td> <td>200 μM with 1.15 nM and 2.87 nM TNFα</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>TNFα reduced expression of GLUT4 and adiponectin, and increased SOCS3 protein levels in these cells. And these effects were prevented.</td> </tr> </table> <p>Cell Invasion Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa and SiHa cells</td> </tr> <tr> <td>Concentration:</td> <td>500 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>72 h</td> </tr> <tr> <td>Result:</td> <td>Significantly reduced hypoxia-elicited cell invasion in both cell models.</td> </tr> </table> <p>Cell Migration Assay^[2]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>HeLa and SiHa cells</td> </tr> </table>	Cell Line:	3T3-L1 adipocytes	Concentration:	200 μM with 1.15 nM and 2.87 nM TNFα	Incubation Time:	72 h	Result:	TNFα reduced expression of GLUT4 and adiponectin, and increased SOCS3 protein levels in these cells. And these effects were prevented.	Cell Line:	HeLa and SiHa cells	Concentration:	500 μM	Incubation Time:	72 h	Result:	Significantly reduced hypoxia-elicited cell invasion in both cell models.	Cell Line:	HeLa and SiHa cells
Cell Line:	3T3-L1 adipocytes																		
Concentration:	200 μM with 1.15 nM and 2.87 nM TNFα																		
Incubation Time:	72 h																		
Result:	TNFα reduced expression of GLUT4 and adiponectin, and increased SOCS3 protein levels in these cells. And these effects were prevented.																		
Cell Line:	HeLa and SiHa cells																		
Concentration:	500 μM																		
Incubation Time:	72 h																		
Result:	Significantly reduced hypoxia-elicited cell invasion in both cell models.																		
Cell Line:	HeLa and SiHa cells																		

Concentration:	500 μ M
Incubation Time:	72 h
Result:	Decreased hypoxia-induced migration from 180 and 240% to 60 and 70% in HeLa and SiHa cells, respectively.
Western Blot Analysis ^[2]	
Cell Line:	HeLa and SiHa cells
Concentration:	500 μ M
Incubation Time:	72 h
Result:	Effectively prevented hypoxia-induced downregulation of E-cadherin and strongly inhibited hypoxia-induced upregulation of α -SMA and vimentin.

In Vivo

β -Aminopropionitrile (BAPN) (100 mg/kg/day; p.o.; 6 weeks) reduces body weight gain and improves the metabolic profile in diet-induced obesity in rats^[1].
 β -Aminopropionitrile monofumarate (1 g/kg/day; p.o.; 4 weeks) induces thoracic aortic dissection in C57BL/6 mice^[3].
MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Male Wistar rats of 150 g, high-fat diet (HFD) model ^[1]
Dosage:	100 mg/kg/day
Administration:	In the drinking water, 6 weeks
Result:	Significantly prevented the rise in body weight in HFD rats, but not in animals that were fed a standard diet. Reduced the increase in the weight of white adipose tissue (both epididymal and lumbar) in obese animals and attenuated their enhanced adiposity. Improved fasted glucose and insulin levels and consequently reduced HOMA index in the HFD group. Improved insulin signalling in adipose tissue from obese animals.
Animal Model:	C57BL/6 mice ^[3]
Dosage:	1 g/kg/day
Administration:	In the drinking water, 4 weeks
Result:	Induce thoracic aortic dissection (TAD) in all mice with 24 h of Ang II infusion. Caused 87% of C57BL/6 mice to develop TAD without Ang II.

REFERENCES

- [1]. Miana M, et al. The lysyl oxidase inhibitor β -aminopropionitrile reduces body weight gain and improves the metabolic profile in diet-induced obesity in rats. *Dis Model Mech.* 2015 Jun;8(6):543-51.
- [2]. Yang X, et al. Inactivation of lysyl oxidase by β -aminopropionitrile inhibits hypoxia-induced invasion and migration of cervical cancer cells. *Oncol Rep.* 2013 Feb;29(2):541-8.
- [3]. Ren W, et al. β -Aminopropionitrile monofumarate induces thoracic aortic dissection in C57BL/6 mice. *Sci Rep.* 2016 Jun 22;6:28149.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA