Proteins



Guanabenz

Cat. No.: HY-12724 CAS No.: 5051-62-7 Molecular Formula: $C_8H_8Cl_2N_4$ Molecular Weight: 231.08

Target: Adrenergic Receptor; Parasite

Pathway: GPCR/G Protein; Neuronal Signaling; Anti-infection

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

Analysis.

Product Data Sheet

BIOLOGICAL ACTIVITY

Description Guanabenz is an orally active α -2-adrenoceptor agonist. Guanabenz has antihypertensive effect and antiparasitic activity. Guanabenz interferes ER stress-signalling and has protective effects in cardiac myocytes. Guanabenz also is used for the

research of high blood pressure^{[1][2][3]}.

IC₅₀ & Target Toxoplasma Toxoplasma

In Vitro Guanabenz (0.5-50 µM, 24 h) is treated with increasing concentrations for 24 hours not affect cell viability^[1].

> Guanabenz (0.5-50 µM, 24 h) alone not affects the UPR targets, neither on mRNA or protein level nor the phosphorylation status of eIF2a. Guanabenz also not induces GADD34 or the constitutively active form $CReP^{[1]}$.

Guanabenz (0.5-50 μ M, 24 h) alone not induces ER stress in neonatal rat cardiomyocytes [1].

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

Cell Viability Assay^[1]

Neonatal rat cardiac myocytes (NRCM)
0.5–50 μM
24 h
Did not affect cell survival.
Neonatal rat cardiac myocytes (NRCM)
0.5–50 μΜ
24 h
Did not affect levels of UPR targets.
Neonatal rat cardiac myocytes (NRCM)

Concentration:	0.5–50 μΜ
Incubation Time:	24 h
Result:	Increased the levels of low panel concentration-dependent UPR targets proteins.

In Vivo

Guanabenz (5 mg/kg/day; i.p.; for 3 weeks) can reproducibly reduce brain cyst burden [2].

Guanabenz (5 mg /kg/d, i.p., oral; 10 mg/kg/d, gavage; for 3 weeks) reverses Toxoplasma-induced hyperactivity in latently infected mice^[2].

Guanabenz (100 and 320 μ g/kg and 1 mg/kg, i.v., over a period of 5 min at intervals of 40 min) reduces sympathetic outflow, heart rate and blood pressure in debuffered cats^[3].

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

Animal Model:	BALB/cJ mice ^[2]
Dosage:	5 mg/kg
Administration:	5 mg/kg/day; i.p. ; for 3 weeks
Result:	Reduced the latent brain cysts in both male and female BALB/cJ mice.
Animal Model:	BALB/cJ mice ^[2]
Dosage:	5 mg/kg; 10 mg/kg
Administration:	5 mg/kg/d, i.p., oral; 10 mg/kg/d, gavage; for 3 weeks
Result:	Reversed parasite-induced hyperactivity to near-baseline levels.
Animal Model:	Cats ^[3]
Dosage:	100 and 320 μg/kg and 1 mg/kg
Administration:	100 and 320 μg/kg and 1 mg/kg, i.v., over a period of 5 min at intervals of 40 min
Result:	Declined markedly blood pressure and nerve activity.

REFERENCES

- $[1]. Christiane \ Neuber, et al.\ Guanabenz\ interferes\ with\ ER\ stress\ and\ exerts\ protective\ effects\ in\ cardiac\ myocytes.\ PLoS\ One.\ 2014\ Jun\ 3;9(6):e98893.$
- [2]. Jennifer Martynowicz, et al. Guanabenz Reverses a Key Behavioral Change Caused by Latent Toxoplasmosis in Mice by Reducing Neuroinflammation. mBio. 2019 Apr 30;10(2):e00381-19.
- [3]. T Baum, et al. Studies on the centrally mediated hypotensive activity of guanabenz. Eur J Pharmacol. 1976 May;37(1):31-44.

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

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