Inhibitors

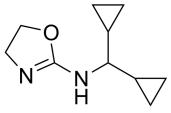
Rilmenidine

Cat. No.: HY-100490 CAS No.: 54187-04-1 Molecular Formula: $C_{10}H_{16}N_{2}O$ Molecular Weight: 180.25

Target: Imidazoline Receptor; Adrenergic Receptor; Apoptosis; Autophagy Pathway: Neuronal Signaling; GPCR/G Protein; Apoptosis; Autophagy

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

Analysis.



Product Data Sheet

BIOLOGICAL ACTIVITY

Description

Rilmenidine, an innovative antihypertensive agent, is an orally active, selective I1 imidazoline receptor agonist. Rilmenidine is an alpha 2-adrenoceptor agonist. Rilmenidine induces autophagy. Rilmenidine acts both centrally by reducing sympathetic overactivity and in the kidney by inhibiting the Na⁺/H⁺ antiport. Rilmenidine modulates proliferation and stimulates the proapoptotic protein Bax thus inducing the perturbation of the mitochondrial pathway and apoptosis in human leukemic K562 cells^{[1][2][3]}.

In Vitro

Rilmenidine provides antihypertensive efficacy comparable with that of diuretics, beta-blockers, calcium channel blockers, and angiotensin-converting enzyme (ACE) inhibitors^[1].

Rilmenidine (25-100 μM; 24 hours) inhibits K562 cell proliferation^[2].

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

Cell Viability Assay^[2]

Cell Line:	K562 cells	
Concentration:	25, 50, 100 μΜ	
Incubation Time:	24 hours	
Result:	Dose-dependently inhibited K562 colony formation.	

In Vivo

Rilmenidine-treated N171-82Q mice (i.p.; 4-times a week) displays significant improved forelimb grip strength and all limbs grip strength from 12 to 22 weeks of age^[3].

Rilmenidine decreases levels of mutant huntingtin^[3].

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

REFERENCES

[1]. Reid JL. Rilmenidine: a clinical overview. Am J Hypertens. 2000;13(6 Pt 2):106S-111S.

[2]. Srdic-Rajic T, et al. Rilmenidine suppresses proliferation and promotes apoptosis via the mitochondrial pathway in human leukemic K562 cells. Eur J Pharm Sci. 2016;81:172-180.

3]. Rose C, et al. Rilmenidine a	ttenuates toxicity of polyglutamine expansions in a mouse mode	of Huntington's disease. Hum Mol Genet. 2010;19(11):2144-2153.
	Caution: Product has not been fully validated for medic	
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