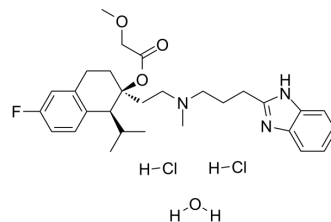


Mibefradil dihydrochloride hydrate

Cat. No.:	HY-15553B
CAS No.:	1049728-52-0
Molecular Formula:	C ₂₉ H ₄₂ Cl ₂ FN ₃ O ₄
Molecular Weight:	586.57
Target:	Calcium Channel
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Mibefradil dihydrochloride hydrate (Ro 40-5967 dihydrochloride hydrate) is an effectively long-acting calcium channel antagonist, used as an antihypertensive agent. Mibefradil dihydrochloride hydrate acts via a higher affinity block for low-voltage-activated (T) than for high-voltage-activated (L) calcium channels ^[1] .
IC₅₀ & Target	Calcium channel ^[1]
In Vitro	Mibefradil inhibits reversibly the T- and L-type currents with IC ₅₀ values of 2.7 and 18.6 μM, respectively ^[2] . Mibefradil (20 μM) reduces the amplitude of excitatory junction potentials (by 37±10 %), slows the rate of repolarisation (by 44±16 %) and causes a significant membrane potential depolarisation (from -83±1 mV to -71±5 mV) ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Br J Pharmacol. 2021 Jan;178(2):346-362.
- J Cell Physiol. 2021 Mar 11.
- Front Pharmacol. 2022 Feb 23;13:816133.
- Front Pharmacol. 23 February 2022.
- Mediat Inflamm. 2020 Nov 10;2020:3691701.

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REFERENCES

- [1]. Martin RL, Lee JH, Cribbs LL, Perez-Reyes E, Hanck DA. Mibefradil block of cloned T-type calcium channels. J Pharmacol Exp Ther. 2000 Oct;295(1):302-8.
- [2]. Mehrke G, et al. The Ca(++)-channel blocker Ro 40-5967 blocks differently T-type and L-type Ca++ channels. J Pharmacol Exp Ther. 1994 Dec;271(3):1483-8.
- [3]. Brain KL, et al. The sources and sequestration of Ca(2+) contributing to neuroeffector Ca(2+) transients in the mouse vas deferens. J Physiol. 2003 Dec 1;553(Pt 2):627-35.

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

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