

Produktinformation



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PRODUCT INFORMATION



Silodosin-d₄ Item No. 30236

CAS Registry No.: 1426173-86-5

Formal Name: (R)-1-(3-hydroxypropyl)-5-(2-((2-(2-

> (2,2,2-trifluoroethoxy)phenoxy)ethyl-1,1,2,2-d₄)amino)propyl)indoline-7-

carboxamide

Synonym: KMD-3213-d₄ MF: $C_{25}H_{28}D_4F_3N_3O_4$

FW: 499.6

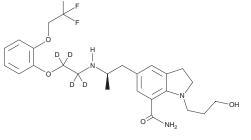
Chemical Purity: ≥98% (Silodosin)

Deuterium

Incorporation: \geq 99% deuterated forms (d₁-d₄); \leq 1% d₀

Supplied as: A solid -20°C Storage: Stability: ≥2 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Laboratory Procedures

Silodosin-d₁ is intended for use as an internal standard for the quantification of silodosin (Item No. 14866) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Silodosin- d_4 is supplied as a solid. A stock solution may be made by dissolving the silodosin- d_4 in the solvent of choice, which should be purged with an inert gas. Silodosin-d₄ is soluble in the organic solvent methanol.

Description

Silodosin is an α_{1A} -adrenergic receptor (α_{1A} -AR) antagonist ($K_i = 0.036$ nM).¹ It is 583- and 56-fold selective for α_{1A} - over α_{1B} - and α_{1D} -ARs, respectively. Silodosin inhibits phenylephrine-induced contraction of isolated rabbit prostate (pA₂ = 10.05) more potently than rabbit or rat aorta (pA₂s = 9.36 and 8.13, respectively).² It inhibits norepinephrine-induced contraction of isolated human prostate tissue when used at concentrations ranging from 0.3 to 10 nM.1 Silodosin (0.01-1,000 µg/kg) inhibits phenylephrineinduced increases in intraurethral pressure in rats.³ Formulations containing silodosin have been used in the treatment of benign prostatic hyperplasia.

References

- 1. Moriyama, N., Akiyama, K., Murata, S., et al. KMD-3213, a novel α_{1A} -adrenoceptor antagonist, potently inhibits the functional α_1 -adrenoceptor in human prostate. Eur. J. Pharmacol. 331(1), 39-42 (1997).
- 2. Yamagishi, R., Akiyama, K., Nakamura, S., et al. Effect of JMD-3213, an α_{1a}-adrenoceptor-selective antagonist, on the contractions of rabbit prostate and rabbit and rat aorta. Eur. J. Pharm. 315(1), 73-79
- 3. Akiyama, K., Hora, M., Tatemichi, S., et al. KMD-3213, a uroselective and long-acting α_{1a} -adrenoceptor antagonist, tested in a novel rat model. J. Pharmacol. Exp. Ther. 291(1), 81-91 (1999).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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