PRODUCT INFORMATION



Doxazosin-d₈ (hydrochloride)

Item No. 30979

CAS Registry No.: 1219803-95-8

Formal Name: [4-(4-amino-6,7-dimethoxy-2-quinazolinyl)-

> 1-piperazinyl-2,2,3,3,5,5,6,6-d_g] (2,3-dihydro-1,4-benzodioxin-2-yl)methanone, monohydrochloride

MF: C₂₃H₁₇D₈N₅O₅ • HCl

FW: 496.0

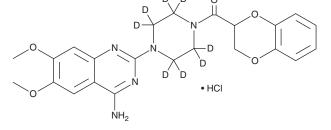
Chemical Purity: ≥98% (Doxazosin)

Deuterium

≥99% deuterated forms (d₁-d₈); ≤1% d₀ Incorporation:

Supplied as: A solid -20°C Storage: Stability: ≥4 years

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



Laboratory Procedures

Doxazosin-d₈ (hydrochloride) is intended for use as an internal standard for the quantification of doxazosin (Item No. 18633) 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).

Doxazosin-d₈ (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the doxazosin-d₈ (hydrochloride) in the solvent of choice, which should be purged with an inert gas. Doxazosin-d₈ (hydrochloride) is soluble in the organic solvent DMSO.

Description

Doxazosin is a non-selective antagonist of α_1 -adrenergic receptors (α_1 -ARs; $K_i s = 3.16$, 1, and 3.98 nM for α_{1A}^{-} , α_{1B}^{-} , and α_{1D}^{-} ARs, respectively). It inhibits norepinephrine-induced contractions in isolated rat aortic rings and human prostate strips with pA_2 values of 8.8 and 8.2, respectively. Doxazosin inhibits phenylephrine-induced increases in blood pressure and prostatic pressure in anesthetized dogs $(pA_2 = 7.5)$ for both). Formulations containing doxazosin have been used in the treatment of benign prostatic hyperplasia and hypertension.

Reference

1. Kenny, B.A., Miller, A.M., Williamson, I.J.R., et al. Evaluation of the pharmacological selectivity profile of α_1 adrenoceptor antagonists at prostatic α_1 adrenoceptors: Binding, functional and in vivo studies. Br. J. Pharmacol. 118(4), 871-878 (1996).

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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