PRODUCT INFORMATION



3-hydroxy Darifenacin

Item No. 27852

CAS Registry No.: 206048-82-0

Formal Name: (3S)-1-[2-(2,3-dihydro-3-hydroxy-

5-benzofuranyl)ethyl]-α,α-

diphenyl-3-pyrrolidineacetamide

Synonym: UK 148993 MF: $C_{28}H_{30}N_2O_3$ FW: 442.6 **Purity:** ≥95%

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

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

Laboratory Procedures

3-hydroxy Darifenacin is supplied as a solid. A stock solution may be made by dissolving the 3-hydroxy darifenacin in the solvent of choice, which should be purged with an inert gas. 3-hydroxy Darifenacin is soluble in organic solvents such as methanol and DMSO, which should be purged with an inert gas.

Description

3-hydroxy Darifenacin is a metabolite of darifenacin (Item No. 14424). 1,2 It is an antagonist of $M_{1,5}$ muscarinic receptors (K,s = 17.78, 79.43, 2.24, 36.31, and 6.17 nM, respectively, in CHO cells).³

References

- 1. Beaumont, K.C., Cussans, N.J., Nichols, D.J., et al. Pharmacokinetics and metabolism of darifenacin in the mouse, rat, dog and man. Xenobiotica 28(1), 63-75 (1998).
- 2. Kerbusch, T., Wählby, U., Milligan, P.A., et al. Population pharmacokinetic modelling of darifenacin and its hydroxylated metabolite using pooled data, incorporating saturable first-pass metabolism, CYP2D6 genotype and formulation-dependent bioavailability. Br. J. Clin. Pharmacol. 56(6), 639-652 (2003).
- 3. Mansfield, K.J., Chandran, J.J., Vaux, K.J., et al. Comparison of receptor binding characteristics of commonly used muscarinic antagonists in human bladder detrusor and mucosa. J. Pharmacol. Exp. Ther. 328(3), 893-899 (2009).

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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