PRODUCT INFORMATION



Cyclopentolate (hydrochloride)

Item No. 23944

CAS Registry No.: 5870-29-1

Formal Name: α-(1-hydroxycyclopentyl)-

benzeneacetic acid, 2-(dimethylamino)

ethyl ester, monohydrochloride

Synonym: (±)-Cyclopentolate MF: C₁₇H₂₅NO₃ • HCl

FW: 327.9 **Purity:** ≥98% 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

Cyclopentolate (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the cyclopentolate (hydrochloride) in the solvent of choice, which should be purged with an inert gas. Cyclopentolate (hydrochloride) is slightly soluble in DMSO and methanol.

Description

Cyclopentolate is an antagonist of muscarinic acetylcholine receptors (Kis = 1.62, 27.5, and 2.63 nM for M₁, M₂, and M₃ receptors, respectively). It inhibits carbachol-induced contraction of isolated human iris sphincter, circular ciliary muscle, and longitudinal ciliary muscle (K_h s = 7.9, 15.8, and 12.5 nM, respectively).² Formulations containing cyclopentolate have been used to induce pupil dilation and to prevent the eye from accommodating for near vision.

References

- 1. Lazareno, S., Buckley, N.J., and Roberts, F.F. Characterization of muscarinic M₄ binding sites in rabbit lung, chicken heart, and NG108-15 cell. Mol. Pharmacol. 38(6), 805-815 (1990).
- 2. Ishikawa, H., DeSantis, L., and Patil, P.N. Selectivity of muscarinic agonists including (±)-aceclidine and antimuscarinics on the human intraocular muscles. J. Ocul. Pharmacol. Ther. 14(4), 363-373 (1998).

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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