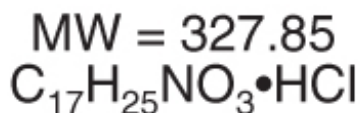
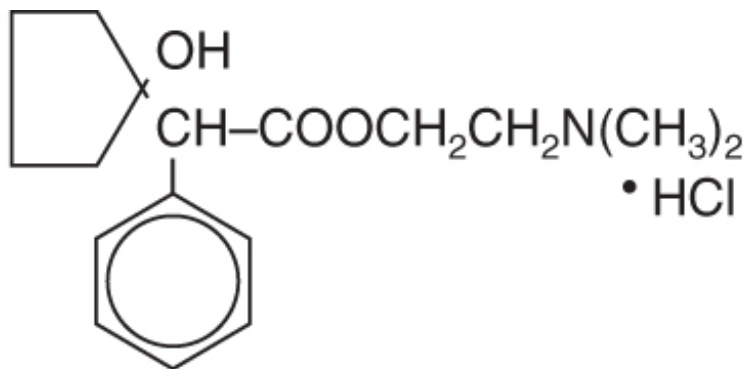


CYCLOPENTOLATE HYDROCHLORIDE- cyclopentolate hydrochloride solution
Sandoz Inc.

Cyclopentolate Hydrochloride Ophthalmic Solution USP

DESCRIPTION

Cyclopentolate Hydrochloride Ophthalmic Solution USP is an anticholinergic prepared as a sterile, borate buffered, solution for topical ocular use. It is supplied in three strengths. The active ingredient is represented by the structural formula:



Established name: Cyclopentolate Hydrochloride

Chemical name: 2-(Dimethylamino)ethyl 1-hydroxy- α -phenylcyclopentaneacetate hydrochloride

Each mL contains: **Active:** cyclopentolate hydrochloride 0.5%, 1% or 2%. **Preservative:** benzalkonium chloride 0.01%. **Inactives:** boric acid, edetate disodium, potassium chloride (except 2% strength), sodium carbonate and/or hydrochloric acid (to adjust pH), purified water. The pH range is between 3.0 and 5.5.

CLINICAL PHARMACOLOGY

This anticholinergic preparation blocks the responses of the sphincter muscle of the iris and the accommodative muscle of the ciliary body to cholinergic stimulation, producing pupillary dilation (mydriasis) and paralysis of accommodation (cycloplegia). It acts rapidly, but has a shorter duration than atropine.

Maximal cycloplegia occurs within 25 to 75 minutes after instillation. Complete recovery of accommodation usually takes 6 to 24 hours. Complete recovery from mydriasis in some individuals may require several days. Heavily pigmented irides may require more doses than lightly pigmented irides.

INDICATIONS AND USAGE

Cyclopentolate hydrochloride is used to produce mydriasis and cycloplegia.

CONTRAINDICATIONS

Should not be used when untreated narrow-angle glaucoma, or untreated anatomically narrow angles are present, or if the patient is hypersensitive to any component of this preparation.

WARNINGS

For topical ophthalmic use only. Not for injection. This preparation may cause CNS disturbances. This is especially true in younger age groups, but may occur at any age, especially with the stronger solutions. Infants are especially prone to CNS and cardiopulmonary side effects from cyclopentolate. To minimize absorption, use only 1 drop of 0.5% Cyclopentolate Hydrochloride Ophthalmic Solution USP per eye, followed by pressure applied over the nasolacrimal sac for two to three minutes. Observe infants closely for at least 30 minutes following instillation.

Mydriatics may produce a transient elevation of intraocular pressure.

PRECAUTIONS

General

The lacrimal sac should be compressed by digital pressure for two to three minutes after instillation to reduce excessive systemic absorption. Caution should be observed when considering use of this medication in the presence of Down's syndrome and in those predisposed to angle-closure glaucoma.

Information for Patients

Do not touch dropper tip to any surface, as this may contaminate the solution. A transient burning sensation may occur upon instillation. Patients should be advised not to drive or engage in other hazardous activities while pupils are dilated. Patients may experience sensitivity to light and should protect eyes in bright illumination during dilation. Parents should be warned not to get this preparation in their child's mouth and to wash their own hands and the child's hands following administration. Feeding intolerance may follow ophthalmic use of this product in infants. It is recommended that feeding be withheld for four (4) hours after examination.

Drug Interactions

Cyclopentolate may interfere with the ocular anti-hypertensive action of carbachol, pilocarpine, or ophthalmic cholinesterase inhibitors.

Carcinogenesis, Mutagenesis, Impairment of Fertility

Studies in animals or humans have not been conducted to evaluate the carcinogenic potential of Cyclopentolate Hydrochloride Ophthalmic Solution USP.

Pregnancy

Pregnancy Category C. Animal reproduction studies have not been conducted with cyclopentolate. It is also not known whether cyclopentolate can cause fetal harm when administered to a pregnant woman or can affect reproduction capacity. Cyclopentolate should be administered to a pregnant woman only if clearly needed.

Nursing Mothers

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when cyclopentolate hydrochloride is administered to a nursing woman.

Pediatric Use

Use of cyclopentolate has been associated with psychotic reactions and behavioral disturbances in pediatric patients. Increased susceptibility to cyclopentolate has been reported in infants, young children, and in children with spastic paralysis or brain damage. These disturbances include ataxia, incoherent speech, restlessness, hallucinations, hyperactivity, seizures, disorientation as to time and place, and failure to recognize people. Feeding intolerance may follow ophthalmic use of this product in infants. It is recommended that feeding be withheld for four (4) hours after examination. Observe infants closely for at least 30 minutes (See WARNINGS).

Geriatric Use

No overall differences in safety or effectiveness between elderly and younger patients.

ADVERSE REACTIONS

Ocular: Increased intraocular pressure, burning, photophobia, blurred vision, irritation, hyperemia, conjunctivitis, blepharoconjunctivitis, punctate keratitis, synechiae have been reported.

Non-ocular: Use of cyclopentolate has been associated with psychotic reactions and behavioral disturbances, usually in children, especially with 2% concentration. These disturbances include ataxia, incoherent speech, restlessness, hallucinations, hyperactivity, seizures, disorientation as to time and place, and failure to recognize people. This drug produces reactions similar to those of other anticholinergic drugs, but the central nervous system manifestations as noted above are more common. Other toxic manifestations of anticholinergic drugs are skin rash, abdominal distention in infants, unusual drowsiness, tachycardia, hyperpyrexia, vasodilation, urinary retention, diminished gastrointestinal motility and decreased secretion in salivary and sweat glands, pharynx, bronchi and nasal passages. Severe manifestations of toxicity include coma, medullary paralysis and death.

OVERDOSAGE

Excessive dosage may produce behavioral disturbances, tachycardia, hyperpyrexia, hypertension, elevated intraocular pressure, vasodilation, urinary retention, diminished gastrointestinal motility and decreased secretion in salivary and sweat glands, pharynx, bronchi and nasal passages. Patients exhibiting signs of overdose should receive supportive care and monitoring.

DOSAGE AND ADMINISTRATION

Adults: Instill one or two drops of 0.5%, 1% or 2% solution in the eye which may be repeated in five to ten minutes if necessary. Complete recovery usually occurs in 24 hours. Complete recovery from mydriasis in some individuals may require several days. **Children:** Instill one or two drops of 0.5%, 1% or 2% solution in the eye which may be repeated five to ten minutes later by a second application of 0.5% or 1% solution if necessary. **Small Infants:** A single instillation of one drop of 0.5% Cyclopentolate Hydrochloride Ophthalmic Solution in the eye. To minimize absorption, apply pressure over the nasolacrimal sac for two to three minutes. Observe infant closely for at least 30 minutes following instillation. Individuals with heavily pigmented irides may require higher strengths.

HOW SUPPLIED

In multiple-dose plastic DROP-TAINER* dispensers:

Cyclopentolate Hydrochloride Ophthalmic Solution USP

0.5%	1%	2%
15 mL NDC 61314-395-01	2 mL NDC 61314-396-01	2 mL NDC 61314-397-01
	5 mL NDC 61314-396-02	5 mL NDC 61314-397-02

15 mL NDC 61314-396-03 15 mL NDC 61314-397-03

Storage: Store at 8° to 25°C (46° to 77°F).

To report SUSPECTED ADVERSE REACTIONS, contact Sandoz Inc., at 1-800-525-8747 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Rx Only

*DROP-TAINER is a registered trademark of Alcon Research, Ltd.

Rev. 10/2015

SANDOZ

a Novartis company

Manufactured by
Alcon Laboratories, Inc.
Fort Worth, Texas 76134 for
Sandoz Inc.
Princeton, NJ 08540
Printed in USA

9012269-0915

PRINCIPAL DISPLAY PANEL

NDC 61314-396-03

Cyclopentolate Hydrochloride Ophthalmic Solution, USP

1%

Rx only

STERILE
15 mL

SANDOZ

a Novartis company

INGREDIENTS: Each mL contains: Active: cyclopentolate hydrochloride 1%. **Preservative:** benzalkonium chloride 0.01%. **Inactives:** boric acid, edetate disodium, potassium chloride, sodium carbonate and/or hydrochloric acid (to adjust pH), purified water. The pH range is between 3.0 and 5.5.

USUAL DOSAGE: Instill one or two drops in the eye which may be repeated in five to ten minutes if necessary.

Read enclosed insert.

PRECAUTION: Do not touch dropper tip to any surface, as this may contaminate the solution.

STORAGE: Store at 8° to 25°C (46° to 77°F).

Manufactured by Alcon Laboratories, Inc.
Fort Worth, Texas 76134 for
Sandoz Inc.
Princeton, NJ 08540

Printed in USA

Rev. 10/2015

9012233-0915

LOT:

EXP.:



NDC 61314-396-03

Cyclopentolate Hydrochloride Ophthalmic Solution, USP
1%

FOR TOPICAL OPHTHALMIC USE ONLY

Rx only

STERILE 15 mL

SANDOZ

INGREDIENTS: Each mL contains: Active: cyclopentolate hydrochloride 1%. **Preservative:** benzalkonium chloride 0.01%. **Inactives:** boric acid, potassium chloride, edetate disodium, sodium carbonate and/or hydrochloric acid (to adjust pH), purified water.

The pH range is between 3.0 and 5.5.

PRECAUTION: Do not touch dropper tip to any surface, as this may contaminate the solution.

Rev. 10/2015

USUAL DOSAGE: Instill one or two drops in the eye which may be repeated in five to ten minutes if necessary. Read enclosed insert.

STORAGE: Store at 8° to 25°C (46° to 77°F).

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Alcon Laboratories, Inc.
Fort Worth, Texas 76134 for
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H14037-0915

LOT/EXP.:

The image shows a detailed product label for Cyclopentolate Hydrochloride Ophthalmic Solution, USP 1%. The label is rectangular with a dashed orange border and a white background. It is divided into several sections. On the left, there is a barcode. The main text is arranged in columns. The top left section lists ingredients: Active (cyclopentolate hydrochloride 1%) and Preservative (benzalkonium chloride 0.01%). The bottom left section lists Inactives (boric acid, potassium chloride, edetate disodium, sodium carbonate and/or hydrochloric acid) and provides pH information (3.0 to 5.5) and a precaution (do not touch dropper tip to any surface). The top center section displays the NDC number (61314-396-03) and the product name in large blue font. Below the product name is a red box with '1%' in white, followed by 'FOR TOPICAL OPHTHALMIC USE ONLY' and 'Rx only STERILE 15 mL'. The bottom center features the Sandoz logo. The top right section provides the usual dosage (instill one or two drops in the eye) and storage instructions (8° to 25°C). The bottom right section lists the manufacturer (Alcon Laboratories, Inc.) and the printer (Sandoz Inc.). On the far right, there is a vertical black bar with 'LOT/EXP.:' and a white box with the product code 'H14037-0915'.

INGREDIENTS: Each mL contains: Active: cyclopentolate hydrochloride 1%. **Preservative:** benzalkonium chloride 0.01%. **Inactives:** boric acid, potassium chloride, edetate disodium, sodium carbonate and/or hydrochloric acid (to adjust pH), purified water.

The pH range is between 3.0 and 5.5. **PRECAUTION:** Do not touch dropper tip to any surface, as this may contaminate the solution.

Rev. 10/2015

NDC 61314-396-03

Cyclopentolate Hydrochloride Ophthalmic Solution, USP

1%

FOR TOPICAL OPHTHALMIC USE ONLY

Rx only STERILE 15 mL

SANDOZ

USUAL DOSAGE: Instill one or two drops in the eye which can be repeated in five to ten minutes if necessary. Read enclosed insert.

STORAGE: Store at 8° to 25°C (46° to 77°F).

Manufactured by Alcon Laboratories, Inc. Fort Worth, Texas 76134 for Sandoz Inc. Princeton, NJ 08540 Printed in USA

LOT/EXP.:

H14037-0915

CYCLOPENTOLATE HYDROCHLORIDE

cyclopentolate hydrochloride solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:61314-396
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CYCLOPENTOLATE HYDROCHLORIDE (UNII: 736I6971TE) (CYCLOPENTOLATE - UNII:I76F4SHP7J)	CYCLOPENTOLATE HYDROCHLORIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM CARBONATE (UNII: 45P3261C7T)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
WATER (UNII: 059QF0KO0R)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61314-396-01	2 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2002	
2	NDC:61314-396-02	5 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2002	
3	NDC:61314-396-03	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/22/2002	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA084110	08/22/2002	

Labeler - Sandoz Inc. (005387188)

Registrant - Alcon Research LLC (007672236)

Establishment

Name	Address	ID/FEI	Business Operations
Alcon Research LLC		007672236	manufacture(61314-396)

