ALTACHLORE SODIUM CHLORIDE HYPERTONICITY- sodium chloride solution/ drops
Altaire Pharmaceuticals Inc.

Altachlore Sodium Chloride Hypertonicity Ophthalmic Solution

iSolutions

ActivEyes

Altachlore5% Solution

Sodium Chloride Hypertonicity

OphthalmicSolution5%

15mL

NDC 59390-183-13

Drug Facts

Activeingredient

Sodium Chloride 5%

Purpose

Hypertonicity Agent

Use

- For the temporary relief of corneal edema.

Warnings

For use in the eyesonly

Save box for complete information

Do not use

- except under the advice and supervision of a doctor.
- if solution changes color or becomes cloudy.

When using this product

- keep tightly closed.
- to avoid contamination, do not touch tip of container to any surface.
- replace cap immediately after each use.
- this product may cause temporary burning and irritation on being instilled into the eye.

Stop use and ask a doctor if

- you experience eye pain.
- changes in your vision occur.
- continued redness or irritation of the eye persists.
- condition worsens or persists.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- instill 1 or 2 drops in the affected eye(s) every 3 to 4 hours, or as directed by a doctor.

Other information

- replace cap after using and keep tightly closed.
- store at 15° to 30°C (59° to 86°F).
- remove contact lenses before using.

Inactive ingredients

Boric Acid, Hypromellose, Methylparaben, Propylene Glycol, Proplyparaben, Sodium Borate, and Water for injection. Sodium Hydroxide and/or Hydrochloric Acid may be added to adjust pH.

Questions?

- 1 (800) 258-2471
- 9am– 5pm EST Monday - Friday

PRINCIPAL DISPLAY PANEL

iSolutionsActivEyesAltachlore5% SolutionSodium ChlorideHypertonicity OphthalmicSolution5%Sterile
15mLNET WT (1/2 FL OZ)

ALTACHLORE SODIUM CHLORIDE HYPERTONICITY

sodium chloride solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59390-183
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59390-183-13	1 in 1 CARTON	09/24/2002	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/24/2002	

Labeler - Altaire Pharmaceuticals Inc. (786790378)

Establishment

Name	Address	ID/FEI	Business Operations
Altaire Pharmaceuticals Inc.		786790378	manufacture(59390-183)

Revised: 12/2023 Altaire Pharmaceuticals Inc.