

SODIUM CHLORIDE HYPERTONICITY OPHTHALMIC SOLUTION- sodium chloride solution
Rugby Laboratories

Drug Facts

Active ingredients

Sodium chloride 5%

Purpose

Hypertonicity agent

Uses

temporary relief of corneal edema

Warnings

Do not use

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

When using this product

- it may cause temporary burning and irritation
- to avoid contamination do not touch tip of container to any surface
- replace cap after use

Stop use and ask a doctor if

- condition worsens or persists for more than 72 hours
- you experience eye pain, changes in vision, continued redness or irritation of the eye

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 or 4 hours, or as directed by a physician.

Other information

- store upright at 15° - 25°C (59° - 77°F)

- keep tightly closed
- serious side effects associated with use of the product may be reported to the phone number provided below

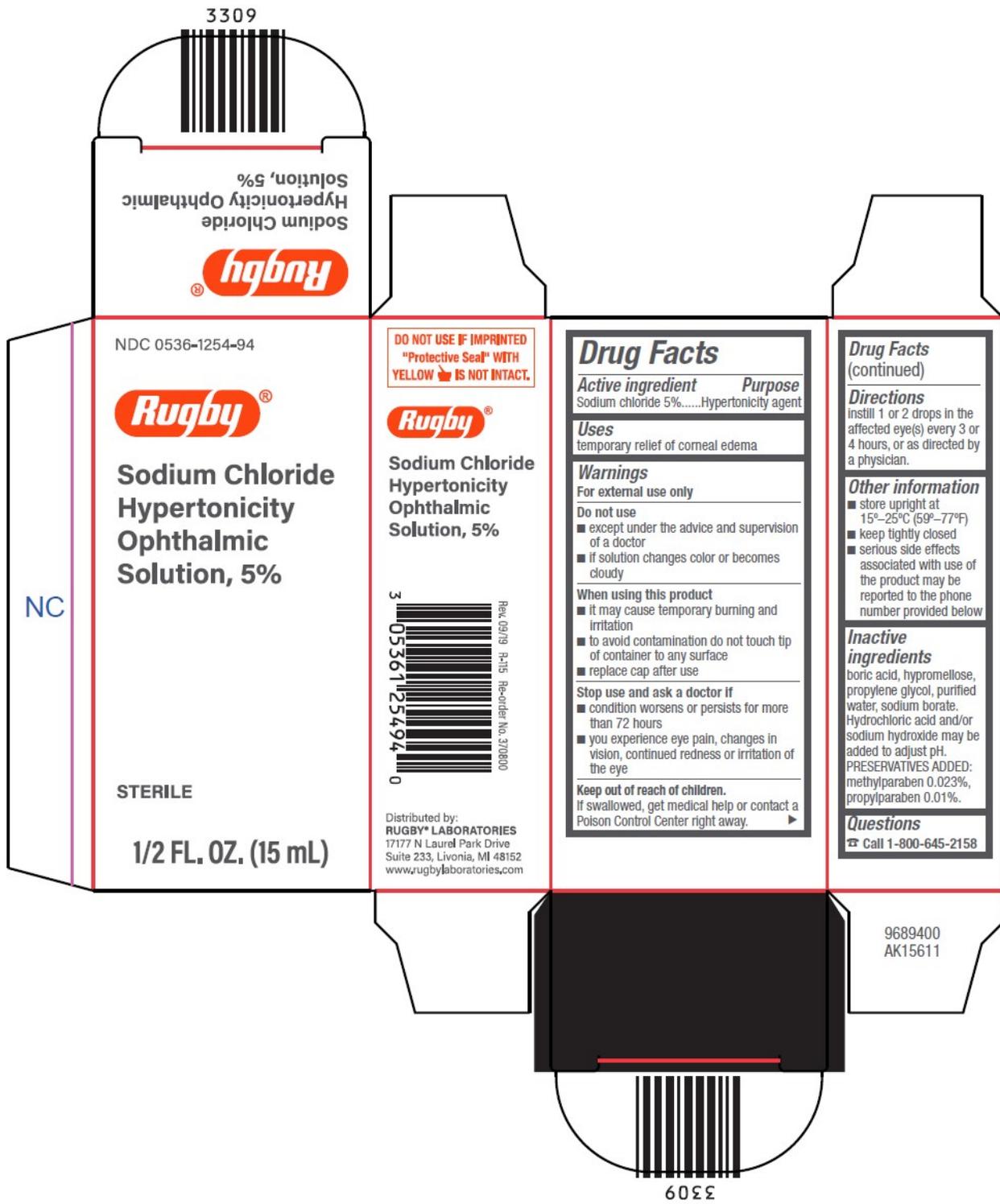
Inactive ingredients

boric acid, hypromellose, propylene glycol, purified water, sodium borate. Hydrochloric acid and/or sodium hydroxide may be added to adjust pH. PRESERVATIVES ADDED: methylparaben 0.023%, propylparaben 0.01%.

Questions?

[phone icon] **Call 1-800-645-2158**

Package/Label Principal Display Panel - carton



Sodium Chloride Hypertonicity Ophthalmic Solution, 5%



NDC 0536-1254-94



Sodium Chloride Hypertonicity Ophthalmic Solution, 5%

NC

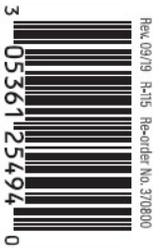
STERILE

1/2 FL. OZ. (15 mL)

DO NOT USE IF IMPRINTED "Protective Seal" WITH YELLOW IS NOT INTACT.



Sodium Chloride Hypertonicity Ophthalmic Solution, 5%



Distributed by:
RUGBY* LABORATORIES
 17177 N Laurel Park Drive
 Suite 233, Livonia, MI 48152
 www.rugbylaboratories.com

Drug Facts

Active ingredient Purpose
 Sodium chloride 5%.....Hypertonicity agent

Uses
 temporary relief of corneal edema

Warnings
 For external use only
 Do not use
 ■ except under the advice and supervision of a doctor
 ■ if solution changes color or becomes cloudy

When using this product
 ■ it may cause temporary burning and irritation
 ■ to avoid contamination do not touch tip of container to any surface
 ■ replace cap after use

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 ■ condition worsens or persists for more than 72 hours
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Drug Facts (continued)

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Questions
 ☎ Call 1-800-645-2158

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NDC 0536-1254-94

Rugby®

Sodium Chloride Hypertonicity Ophthalmic Solution, 5%

STERILE

1/2 FL. OZ. (15 mL)

SODIUM CHLORIDE HYPERTONICITY OPHTHALMIC SOLUTION

sodium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0536-1254
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 2208 (100 MPA.S) (UNII: B1QE5P712K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0536-1254-94	1 in 1 CARTON	09/17/2020	
1		15 mL in 1 BOTTLE; 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	09/17/2020	

Labeler - Rugby Laboratories (079246066)

Establishment

Name	Address	ID/FEI	Business Operations
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Bausch & Lomb Incorporated		079587625	manufacture(0536-1254)
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Revised: 9/2024

Rugby Laboratories