SODIUM CHLORIDE HYPERTONICITY OPHTHALMIC- sodium chloride ointment
Rugby Laboratories

Drug Facts

Active ingredient

Sodium chloride 50 mg (5%)

Purpose

Hypertonicity agent

Uses

temporary relief of corneal edema

Warnings

For external use only

Do not useexcept under the advice and supervision of a doctor

When using this product

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

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

- pull down the lower lid of the affected eye
- apply a small amount (1/4 inch) of ointment to the inside of eyelid
- apply every 3 or 4 hours or as directed by a doctor

Other information

- store at 15° - 25°C (59° - 77°F)
- keep tightly closed
- **DO NOT FREEZE**

- see crimp of tube or carton for Lot Number and Expiration Date
- do not use if difficult to dispense or visible particles are seen in the product
- serious side effects associated with use of the product may be reported to the phone number below

Inactive ingredients

lanolin, mineral oil, purified water, white petrolatum

***Questions*[phone icon]**

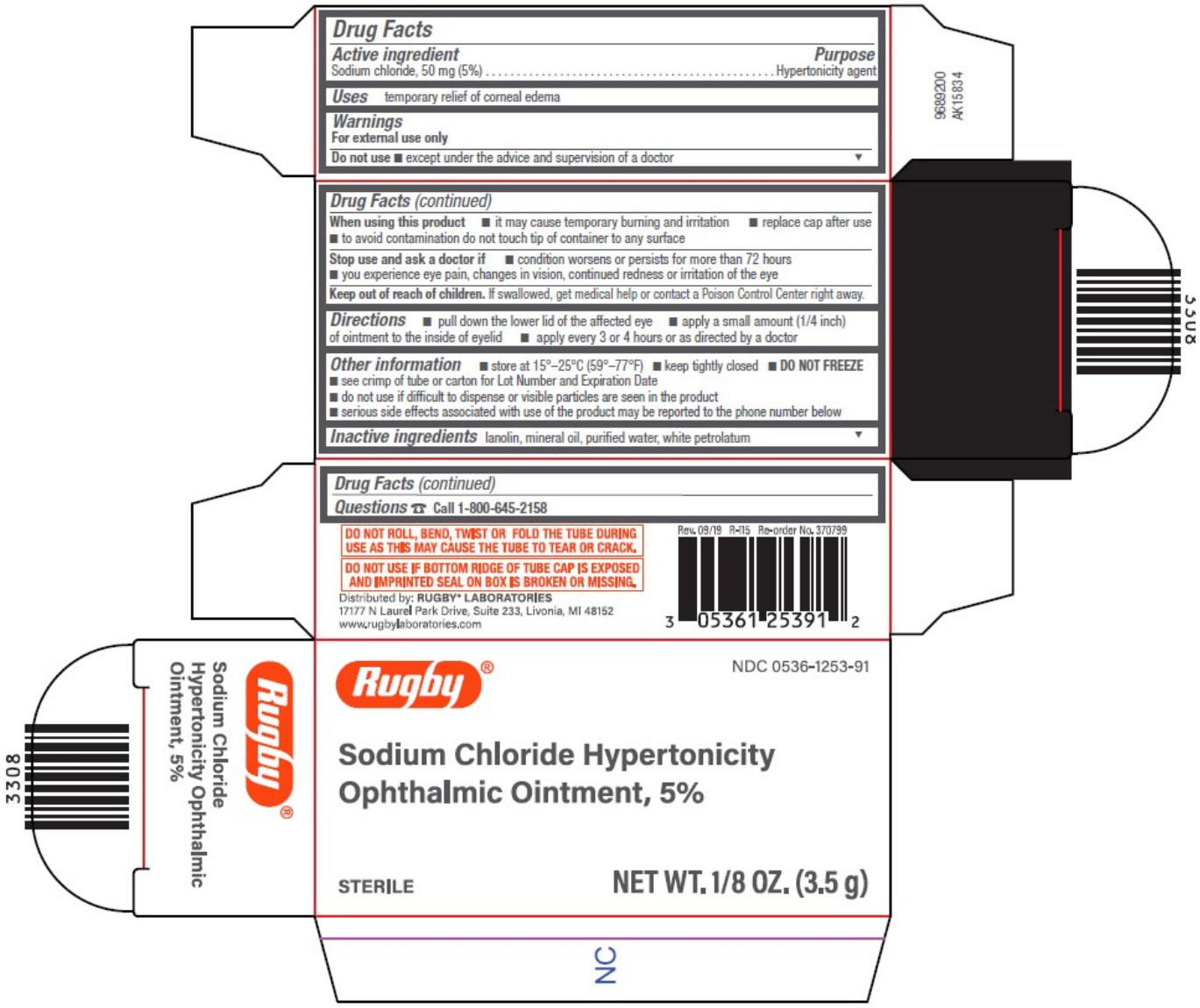
Call 1-800-645-2158

Package/Label Principal Display Panel Carton

***Rugby*®**

NDC 0536-1253-91

Sodium Chloride Hypertonicity
Ophthalmic Ointment, 5%
1. **STERILE NET WT. 1/8 OZ. (3.5 G)**



SODIUM CHLORIDE HYPERTONICITY OPHTHALMIC

sodium chloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0536-1253
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 g

Inactive Ingredients	
Ingredient Name	**Strength**
LANOLIN (UNII: 7EV65EAW6H)	

MINERAL OIL (UNII: T5L8T28FGP)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0536-1253-91	1 in 1 CARTON	11/06/2020	
1		3.5 g in 1 TUBE; 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	11/06/2020	

Labeler - Rugby Laboratories (079246066)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	manufacture(0536-1253)

Revised: 9/2024 Rugby Laboratories