SODIUM CHLORIDE HYPERTONICITY- sodium chloride ointment
CVS PHARMACY

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 use
■except 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 1-866-767-9161

Package/Label Principal Display Panel - Carton

[heart icon] **CVS**Health ™

Compare to the active
ingredient in Muro 128 ®*

NDC 69842-285-35

Sodium Chloride
Hypertonicity
ophthalmic ointment, 5%
Temporary relief of corneal edema

NET WT 0.125 OZ (3.5 g)

STERILE

SODIUM CHLORIDE HYPERTONICITY

sodium chloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-285
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:69842-285-35	1 in 1 CARTON	05/01/2020	
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	05/01/2020	

Labeler - CVS PHARMACY (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	manufacture(69842-285)

Revised: 9/2024 CVS PHARMACY