

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] **CVSHealth**™

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: 059QF0K00R)				
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: 1/2026

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