#### SODIUM CHLORIDE HYPERTONICITY- sodium chloride solution CVS Pharmacy

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# **Drug Facts**

## Active ingredient

Sodium chloride 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
- 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

# 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 1-866-767-9161

# Package/Label Principal Display Panel

[heart icon] **CVS** Health ™

Compare to the active ingredient in Muro 128 <sup>®</sup>\* NDC 69842-284-15

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

Actual Size Bottle on Side Panel

# STERILE

15 mL (0.5 FL OZ)



## SODIUM CHLORIDE HYPERTONICITY

sodium chloride solution

#### **Product Information**

**Product Type** 

HUMAN OTC DRUG

Item Code (Source)

NDC:69842-284

Active Ingredient/Active Moiety							
		Ingredient Name	<b>Basis of Strengt</b>	h Strength			
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37) SODIUM CHLO			7) SODIUM CHLORIDE	50 mg in 1 m			
In	active Ingre	dients					
		Strength					
BC							
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)							
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)							
w							
SODIUM BORATE (UNII: 91MBZ8H3QO)							
HYDROCHLORIC ACID (UNII: QTT17582CB)							
SODIUM HYDROXIDE (UNII: 55X04QC32I)							
50							
		(UNII: A2I8C7HI9T)					
ME	THYLPARABEN						
ME	THYLPARABEN	(UNII: A2I8C7HI9T)					
ME	THYLPARABEN	(UNII: A2I8C7HI9T)					
ME PR	THYLPARABEN	(UNII: A2I8C7HI9T)					
ME PR	THYLPARABEN OPYLPARABEN	(UNII: A2I8C7HI9T) (UNII: Z8IX2SC1OH)	rketing Start Ma Date	arketing End Date			
ме PR <b>Ра</b> #	ackaging Item Code	(UNII: A2I8C7HI9T) (UNII: Z8IX2SC1OH)	Date	-			
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ME PR <b>P</b> a #	OPYLPARABEN OPYLPARABEN ACKaging Item Code NDC:69842-284-	(UNII: A2I8C7HI9T) (UNII: Z8IX2SC1OH) Package Description Ma 1 in 1 CARTON 05/01 15 mL in 1 BOTTLE; Type 0: Not a Combination	Date	-			
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ME PR Pa 1	ACKAGING Item Code NDC:69842-284- 15	(UNII: A2I8C7HI9T)   (UNII: Z8IX2SC10H)   Package Description   1 in 1 CARTON   05/01   15 mL in 1 BOTTLE; Type 0: Not a Combination Product   Information	<b>Date</b> /2021	-			

# Labeler - CVS Pharmacy (062312574)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Bausch & Lomb Incorporated		079587625	manufacture(69842-284)				

Revised: 9/2024

**CVS** Pharmacy