MURO 128- sodium chloride solution
Bausch & Lomb Incorporated

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Muro 128

Drug Facts

Active ingredient

Sodium chloride 2%

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 doctor.

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, methylparaben, propylene glycol, propylparaben, purified water, sodium borate. Hydrochloric acid and/or sodium hydroxide may be added to adjust pH.

Questions or comments?

[telephone icon] **Call 1-800-553-5340**

Marketed by:

Bausch & Lomb Americas Inc.

Bridgewater, NJ 08807 USA

9756801

AB15511

Package/Label Principal Display Panel

BAUSCH + LOMB

NDC 24208-276-15

Muro 128®

sodium chloride hypertonicity ophthalmic solution, 2%

SOLUTION

2%

Temporary Relief of Corneal Edema

STERILE

1/2 FL OZ (15mL)

MURO 128

sodium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-276
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	20 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:24208-276-15	1 in 1 CARTON	01/01/2011	
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 final	part349	01/01/2011	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	MANUFACTURE(24208-276)

Revised: 5/2022

Bausch & Lomb Incorporated