MURO 128- sodium chloride ointment
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 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 provided below

Inactive ingredients

lanolin, mineral oil, purified water, white petrolatum

Questions or comments?

Call 1-800-553-5340

Package/Label Principal Display Panel

BAUSCH +LOMB

NDC 24208-385-56

Muro 128®

sodium chloride hypertonicity

ophthalmic ointment, 5%

OINTMENT

5%|

Temporary Relief

of Corneal Edema

TWIN

PACK

STERILE

NET WT. 1/4 OZ. (7 g)

TWO 1/8 OZ (3.5 g) Tubes

9758101
AB15899

MURO 128

sodium chloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24208-385
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)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24208-385-55	1 in 1 CARTON	01/01/2011	
1		3.5 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:24208-385-56	2 in 1 CARTON	01/01/2011	
2		3.5 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:24208-385-01	1 in 1 CARTON	01/01/2011	
3		1 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 final	part349	01/01/2011	

Labeler - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		079587625	MANUFACTURE(24208-385)

Revised: 4/2022 Bausch & Lomb Incorporated