WALGREENS SODIUM CHLORIDE OPHTHALMIC 5 PERCENT HYPERTONICITY
EYE- sodium chloride ointment
Walgreen Company

Walgreens 5% Hypertonicity Eye Ointment (PLD)

Walgreens Sodium Chloride Ophthalmic

Ointment 5% Hypertonicity Eye Ointment 3.5g

NDC 0363-7500-50

Drug Facts

Active ingredient

Sodium Chloride, 50 mg (5%)

Purpose

Hypertonicity Agent

Use

• For the temporary relief of corneal edema.

Warnings

• **For use in the eyesonly.**

• Retain outer carton for full product drug facts.

Do not use

this product except under the advice and supervision of a doctor.

When using this product

• avoid contamination, do not touch tip of container to any surface.

• replace cap after use.

• this product may cause temporary burning and irritation on being instilled into the eye.

Stop use and ask a doctorif

• you experience eye pain, changes in vision, continued redness or irritation of the eye.

• condition worsens or persists for more than 72 hours.

KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• pull down lower lid of the affected eye(s) and apply a small amount (1/4 inch) to the inside of the eyelid and apply every 3-4 hours, or as directed by a doctor.

Other information

• Store at room temperature 15°- 30°C (59°- 86°F).

• Do not freeze.

• Keep tightly closed.

• See crimp of tube or box for lot number and expiration date.

Inactive ingredients

lanolin alcohol, mineral oil, water for injection and white petrolatum.

Questionsor comments?

•1- 800-925-4733

PRINCIPAL DISPLAY PANEL

NDC 0363-7500-50

Sodium Chloride

Ophthalmic

Ointment, 5%

Hypertonicity

Eye Ointment

NET WT 0.125 OZ (3.5 g)

Void Aqueous

Walgreens

DO NOT USE IF BOTTOM RIDGE OF TUBE IS EXPOSED, OR IF CAP OR TUBE IS DAMAGED

Drug Facts

Active ingredient	Purpose
Sodium chloride, (5%)	Hypertonicity agent

Use
- For the temporary relief of corneal edema.

Warnings
- **For use in the eyes only.**
- Retain outer carton for full product drug facts.

Do not use this product except under the advice and supervision of a doctor.

When using this product
- Avoid contamination, do not touch tip of container to any surface.
- Replace cap after use.
- This product may cause temporary burning and irritation on being instilled into the eye.

Stop use and ask a doctor if
- You experience eye pain, changes in vision, continued redness or irritation of the eye.
- Condition worsens or persists for more than 72 hours.

Keep this and all drugs out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- Pull down lower lid of the affected eye(s) and apply a small amount (1/4 inch) to the inside of the eyelid and apply every 3-4 hours, or as directed by a doctor.

Drug Facts (continued)

Other information
- Store at room temperature 15°-30°C (59°-86°F).
- Do not freeze.
- Keep tightly closed.
- See crimp of tube or box for lot number and expiration date.

Inactive ingredients
lanolin alcohol, mineral oil, purified water and white petrolatum.

Questions or comments?
1-800-925-4733

DISTRIBUTED BY: WAL-GREEN CO., 200 WILMOT RD. DEERFIELD, IL 60015 **100% SATISFACTION GUARANTEED** walgreens.com ©2020 Walgreen Co.

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Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey

[1] This product is not manufactured or distributed by Bausch & Lomb, owner of the registered trademark Muro 128® Hypertonicity Ophthalmic Ointment, 5%.

Walgreens

Compare to Muro 128® Hypertonicity Ophthalmic Ointment, 5% active ingredient[1]

NDC 0363-7500-50

Sodium Chloride Ophthalmic Ointment, 5% Hypertonicity Eye Ointment

- For temporary relief of corneal edema
- Sterile

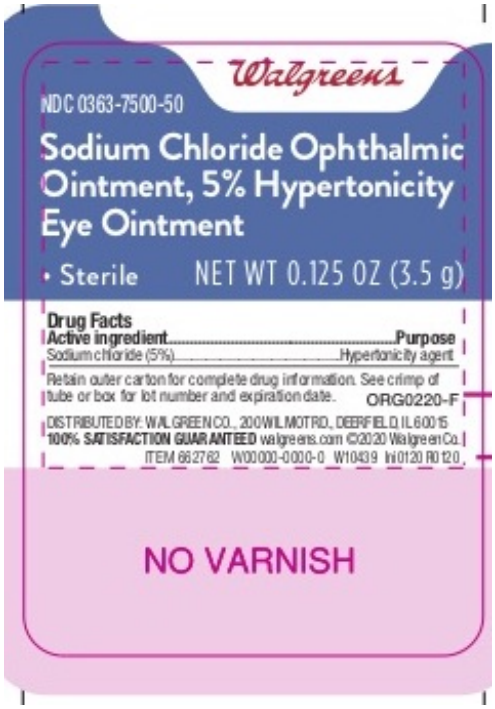
NET WT 0.125 OZ (3.5 g)

ACTUAL SIZE

W10439

ORG0220-F
Ini0120
R0120

Void Aqueous



WALGREENS SODIUM CHLORIDE OPHTHALMIC 5 PERCENT HYPERTONICITY EYE

sodium chloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-7500
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	50 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
MINERAL OIL (UNII: T5L8T28FGP)	
WATER (UNII: 059QF0KO0R)	
PETROLATUM (UNII: 4T6H12BN9U)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0363-7500-50	1 in 1 CARTON	04/01/2019	
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	04/01/2019	

Labeler - Walgreen Company (008965063)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Revised: 12/2024 Walgreen Company