

SODIUM CHLORIDE- sodium chloride ointment

Walgreens

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.

Drug Facts

Active ingredient

Sodium Chloride 5%

Purpose

Hypertonicity Agent

Use

For temporary relief of corneal edema.

Warnings

- do not use this product except under the advice and supervision of a doctor
- do not use if cap skirt is dislodged from tube
- to avoid contamination, do not touch tip of container to any surface
- replace cap after using
- may cause temporary burning and irritation upon application into the eye

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

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

Directions

Apply small amount (one-fourth inch) to the inside of affected eye(s) every 3 to 4 hours, or as directed by a doctor.

Other Information

- store at 20° to 25° C (68° to 77°F) [see USP Controlled Room Temperature].
- store away from heat

- protect from freezing
- keep tightly closed
- see crimp for Lot Number and Expiration Date
- **Retain this carton for future reference**

Inactive ingredients

- Mineral Oil, Modified Lanolin, Water for Injection and White Petrolatum.

Questions or comments?

1-800-932-5676

Principal Display Panel Text for Container Label:

STERILE Well at



Sodium Chloride Walgreens

Ophthalmic Ointment NDC 0363-9050-00

USP, 5% NET WT 3.5 g (1/8 oz.)

Hypertonicity Eye Ointment

FOR OPHTHALMIC USE ONLY

<p>STERILE Sodium Chloride Ophthalmic Ointment USP, 5% Hypertonicity Eye Ointment FOR OPHTHALMIC USE ONLY</p>		<p>NDC 0363-9050-00</p>
<p>NET WT 3.5 g (1/8 oz.)</p>		
<p>READ OUTER CARTON FOR INFORMATION BEFORE USING. ACTIVE INGREDIENT: Sodium Chloride 5% (50 mg/g) INACTIVES: Mineral Oil, Modified Lanolin, Water for Injection and White Petrolatum. INDICATIONS: For temporary relief of corneal edema. DIRECTIONS: Apply small amount (one-fourth inch) to the inside of affected eye (s) every 3 to 4 hours, or as directed by a doctor. See crimp for Lot Number and Expiration Date. STORAGE: Store at 20° to 25° C (68° to 77° F) [see USP Controlled Room Temperature]. KEEP TIGHTLY CLOSED. KEEP OUT OF REACH OF CHILDREN. W10384-0816-O</p>		
<p style="border: 1px solid black; display: inline-block; padding: 2px;">DO NOT USE IF BOTTOM RIDGE OF TUBE CAP IS EXPOSED</p>		
<p>DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015 100% SATISFACTION GUARANTEED walgreens.com ©2013 Walgreen Co.</p>	<p>ITEM 927924 ORG0113-F2</p>  <p>(01) 00303639050003</p>	<p style="writing-mode: vertical-rl; transform: rotate(180deg);"> LT-1065-2 WAGKOAL Rev. 03/17 </p>

Principal Display Panel Text for Carton Label:

STERILE Well at

Walgreens NDC 0363-9050-00

WALGREENS PHARMACIST RECOMMENDED[†]

Sodium Chloride

Ophthalmic Ointment

USP, 5%

Hypertonicity Eye Ointment

Compare to Muro 128[®]

Ointment active ingredient[#]

NET WT 3.5 g (1/8 OZ)

Drug Facts (continued)

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Directions Apply small amount (one-fourth inch) to the inside of affected eye(s) every 3 to 4 hours, or as directed by a doctor.

Drug Facts (continued)

Other information

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Inactive ingredients Mineral Oil, Modified Lanolin, Water for Injection and White Petrolatum.

Questions or comments? 1-800-932-5676

ORIG 114-F-PO1

WAGK0AC Rev. 08/16

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ITEM 927924 WT 0160-0816-L

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HOW TO SUBSCRIBE: 800.421.0000 or walgreens.com ©2014 Walgreens Co.
MADE IN U.S.A.

STERILE

Sodium Chloride Ophthalmic Ointment USP, 5%

Hypertonicity Eye Ointment

Well at Walgreens

WALGREENS PHARMACIST RECOMMENDED[†]

Compare to Muro 128[®] Ointment active ingredient^{††}

NET WT 3.5 g (1/8 OZ)

Drug Facts

Active ingredient	Purpose
Sodium Chloride 5%	Hypertonicity Agent

Use For temporary relief of corneal edema.

FOR TOPICAL OPHTHALMIC USE ONLY

[†]Walgreens Pharmacist Survey Study, November 2012.

^{††}This product is not manufactured or distributed by Bausch & Lomb Pharmaceuticals, Inc., owner of the registered trademark Muro 128[®].

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SODIUM CHLORIDE

sodium chloride ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9050
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
Mineral Oil (UNII: T5L8T28FGP)	
Lanolin (UNII: 7EV65EAW6H)	
Water (UNII: 059QF0KO0R)	
Petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9050-00	1 in 1 CARTON	02/12/2013	
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 final	part349	02/12/2013	

Labeler - Walgreens (008965063)

Registrant - Akorn Operating Company LLC (117693100)

Establishment

Name	Address	ID/FEI	Business Operations
Akorn		117696840	MANUFACTURE(0363-9050) , ANALYSIS(0363-9050) , STERILIZE(0363-9050) , PACK(0363-9050) , LABEL(0363-9050)

