GENTEAL TEARS MODERATE PRESERVATIVE FREE- dextran 70, hypromellose 2910 solution/ drops

Alcon Laboratories, Inc.

Drug Facts

Active Ingredients	Purpose
Dextran 70 0.1%	Lubricant
Hypromellose 2910 0.3%	Lubricant

Uses

 temporary relief of discomfort due to minor irritations of the eye, or to exposure to wind or sun

Warnings

For external use only

Do not use

- if this solution changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product

- to avoid contamination, do not touch tip of container to any surface
- do not reuse
- once opened, discard

Stop use and ask a doctor if

you experience any of the following:

- eye pain
- changes in vision
- continued redness or irritation of the eye
- 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

put 1 or 2 drops in the affected eye(s) as needed.

Other information

■ store are room temperature 15° – 25° C (59° – 77° F)

Inactive ingredients

potassium chloride, purified water, sodium borate, sodium chloride. May contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions?

In the U.S. call 1 (800) 757-9195 (Mon-Fri 9AM-5PM CST) alcon.medinfo@alcon.com

PRINCIPAL DISPLAY PANEL PRESERVATIVE FREE

Moderate DRY EYE SYMPTOM RELIEF

GenTeal® Tears

LUBRICANT EYE DROPS

LIQUID DROPS

Fast, Soothing Relief Preservative Free Formula

Previously TEARS NATURALE® FREE LUBRICANT EYE DROPS

STERILE 36 Single-Use Vials 0.9 mL (0.03 FL OZ) Each

Alcon

GenTeal[®] Tears Lubricant Eye Drops Preservative Free provide lasting relief for moderate dry eye with the convenience of a vial.

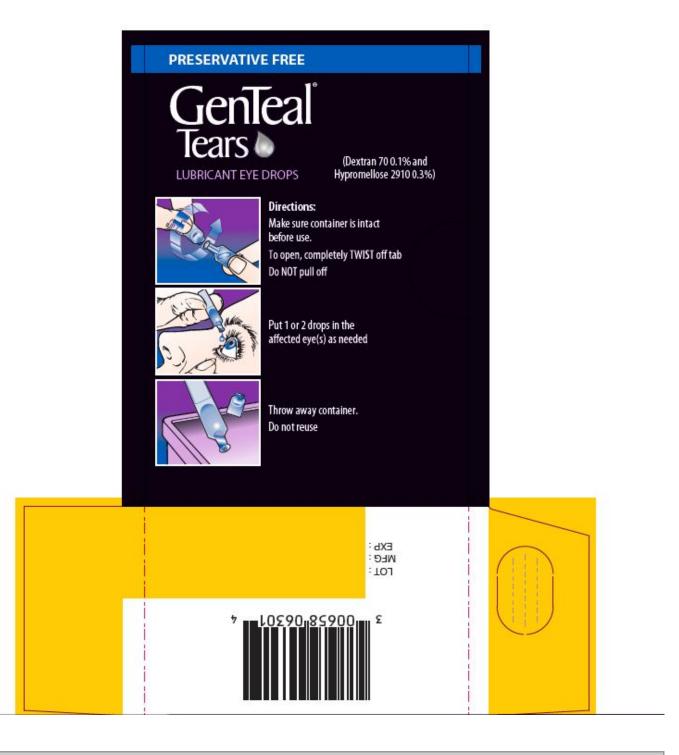
TAMPER EVIDENTY:

Do not use if vial is damaged or opened.

Distributed by: Alcon Laboratories, Inc., 6201 South Freeway Fort Worth, Texas 76134 Country of Origin: France

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GENTEAL TEARS MODERATE PRESERVATIVE FREE

dextran 70, hypromellose 2910 solution/ drops

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Sou	rce)	NDC:0065	-8063
Route of Administration	OPHTHALMIC				
Active Ingredient/Active Moiety					
Ingredient Name Basis of Strength Str			Strength		
Hypromellose 2910 (4000 Mpa.s) (UNII: RN3152OP35) (Hypromellose 2910			Hypromellose	2910	3 mg

(4000 Mpa.s) - UNII:RN3152OP35)	(4000 Mpa.s)	in 1 mL
Dextran 70 (UNII: 7SA290YK68) (Dextran 70 - UNII:7SA290YK68)	Dextran 70	1 mg in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
Potassium Chloride (UNII: 660YQ98I10)				
Water (UNII: 059QF0KO0R)				
Sodium Borate (UNII: 91MBZ8H3QO)				
Hydrochloric Acid (UNII: QTT17582CB)				
Sodium Hydroxide (UNII: 55X04QC32I)				
Sodium Chloride (UNII: 451W47IQ8X)				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0065- 8063-01	36 in 1 CARTON	07/03/2018		
1		.9 mL in 1 VIAL, SINGLE-DOSE; 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	07/03/2018	

Labeler - Alcon Laboratories, Inc. (008018525)

Establishment				
Name	Address	ID/FEI	Business Operations	
Kaysersberg Pharmaceuticals		267486052	manufacture(0065-8063)	

Revised: 12/2023 Alcon Laboratories, Inc.