ALCON TEARS LUBRICANT EYE DROPS- hypromellose 2910 solution/ drops
Alcon Laboratories, Inc.

Drug Facts

Active ingredients	***Purpose***
Hypromellose 2910 0.5%	Lubricant

Uses

- for the temporary relief of burning and irritation due to dryness of the eye
- for use as a protectant against further irritation

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

- remove contact lenses before using
- to avoid contamination, do not touch tip of container to any surface
- replace cap after each use

Stop use and ask a doctor if

you experience any of the following:

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

- instill 1 or 2 drops in the affected eye(s) as needed

Other information

- store at room temperature

Inactive ingredients

benzalkonium chloride 0.01% as preservative, dibasic sodium phosphate, monobasic sodium phosphate, purified water, sodium chloride, sodium citrate

Questions?

In the U.S. call 1-800-757-9195

PRINCIPAL DISPLAY PANEL

Alcon Tears

LUBRICANT
Eye Drops

For Relief of Dry Eye Symptoms

STERILE

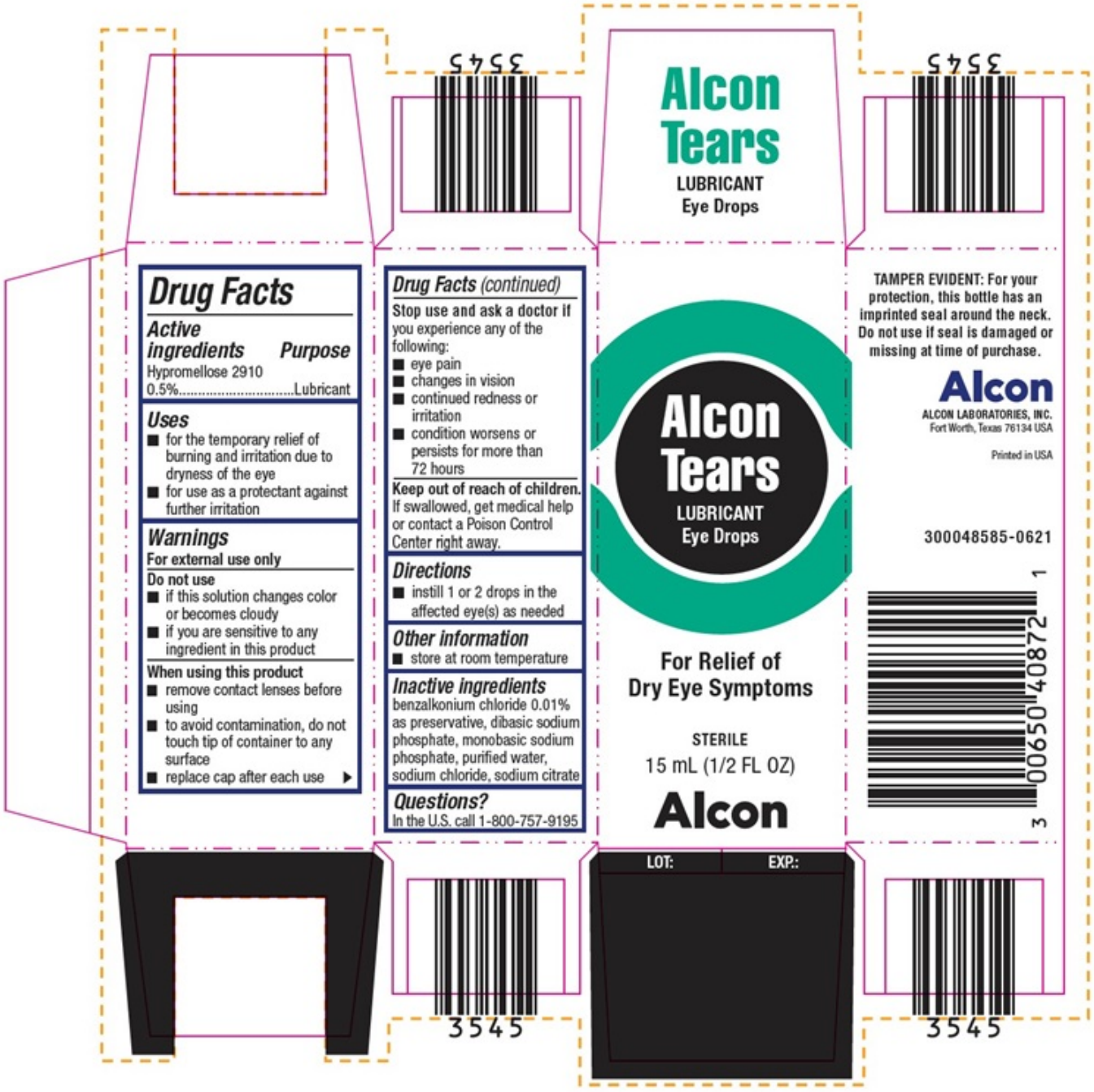
15 mL (1/2 FL OZ)

Alcon

TAMPER EVIDENT: For your protection, this bottle has an imprinted seal around the neck.
Do not use if seal is damaged or missing at time of purchase.

Alcon
ALCON LABORATORIES, INC.
Fort Worth, Texas 76134 USA

Printed in USA



ALCON TEARS LUBRICANT EYE DROPS

hypromellose 2910 solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0065-0408
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Hypromellose 2910 (4000 Mpa.S) (UNII: RN3152OP35) (Hypromellose 2910 (4000 Mpa.S) - UNII:RN3152OP35)	Hypromellose 2910 (4000 Mpa.S)	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7)	
Sodium Phosphate, Dibasic, Unspecified Form (UNII: GR686LBA74)	
Sodium Phosphate, Monobasic, Unspecified Form (UNII: 3980JIH2SW)	
Water (UNII: 059QF0KO0R)	
Sodium Chloride (UNII: 451W47IQ8X)	
Sodium Citrate, Unspecified Form (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0065-0408-72	1 in 1 CARTON	03/01/2022	
1		15 mL in 1 BOTTLE, DROPPER; 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	03/01/2022	

Labeler - Alcon Laboratories, Inc. (008018525)

Establishment

Name	Address	ID/FEI	Business Operations
Alcon Research LLC		007672236	manufacture(0065-0408)

Revised: 12/2023 Alcon Laboratories, Inc.