ISOPTO TEARS- hypromellose 2910 solution
Alcon Laboratories, Inc.

Drug Facts

OTC - ACTIVE INGREDIENT SECTION

Hypromellose 2910 0.5%

OTC - PURPOSE SECTION

Lubricant

INDICATIONS & USAGE SECTION

Uses

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

WARNINGS SECTION

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.

OTC - WHEN USING SECTION

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.

OTC - STOP USE SECTION

Stop use and ask a doctor if

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

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

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

DOSAGE & ADMINISTRATION SECTION

Directions

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

OTHER INFORMATION

- Store at room temperature.

INACTIVE INGREDIENT SECTION

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

OTC - QUESTIONS SECTION

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

PRINCIPAL DISPLAY PANEL

Alcon®

Isopto® Tears
LUBRICANT
Eye Drops
Hypromellose (Ophthalmic Solution)

For Relief of Dry Eyes

STERILE
15 mL (1/2 FL OZ)

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.

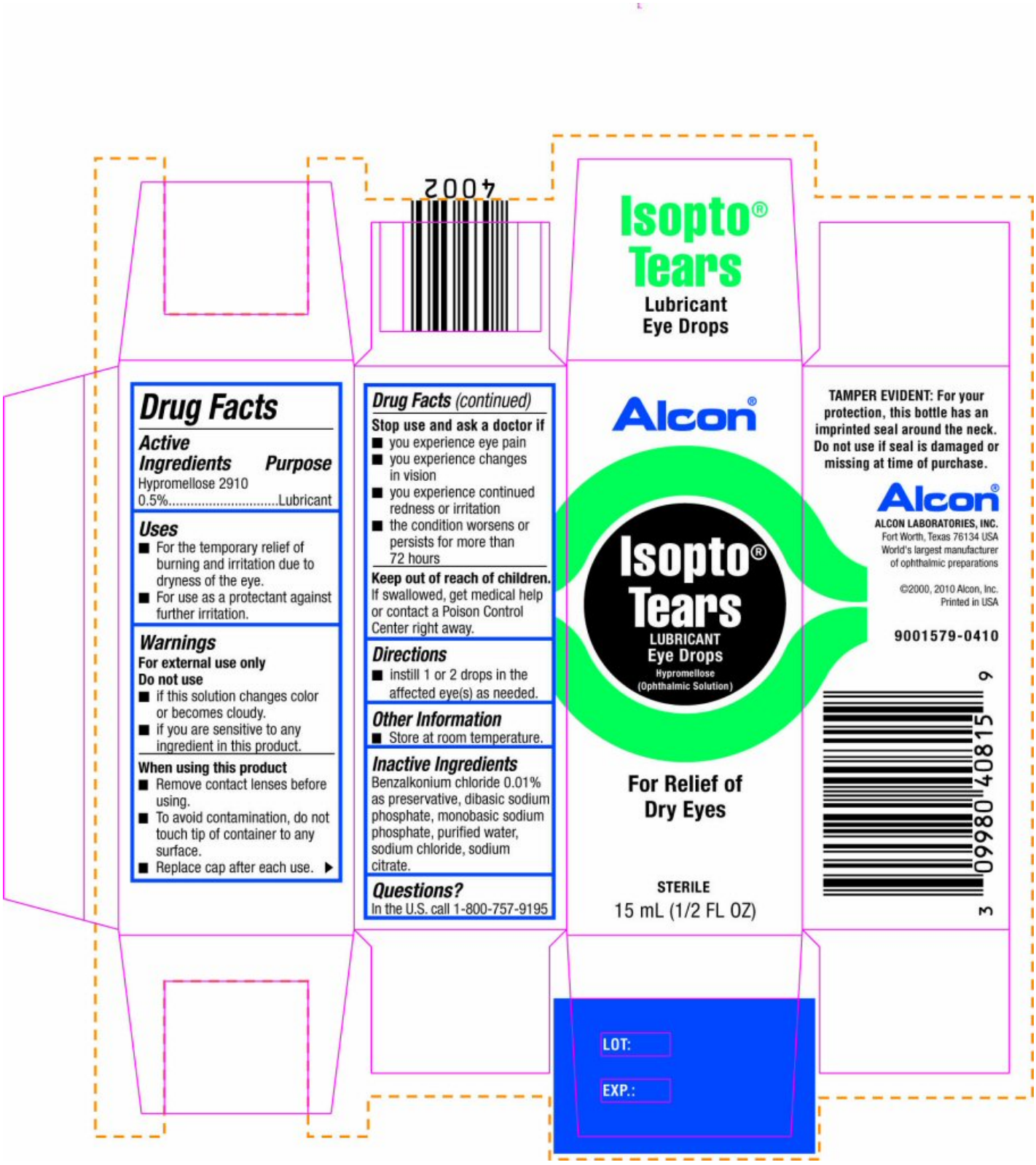
Alcon®

ALCON LABORATORIES, INC.
Fort Worth, Texas 76134 USA

World's largest manufacturer of ophthalmic preparations

Printed in USA

9001579-0410



ISOPTO TEARS

hypromellose 2910 solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0998-0408
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 (UNII: GR686LBA74)	
Sodium Phosphate, Monobasic (UNII: 3980JIH2SW)	
Water (UNII: 059QF0KO0R)	
Sodium Chloride (UNII: 451W47IQ8X)	
Sodium Citrate (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0998-0408-15	15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/30/1990	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	09/30/1990	12/31/2024

Labeler - Alcon Laboratories, Inc. (008018525)

Registrant - Alcon Laboratories, Inc. (008018525)

Establishment

Name	Address	ID/FEI	Business Operations
Alcon Research Ltd		007672236	MANUFACTURE(0998-0408)

Revised: 12/2023 Alcon Laboratories, Inc.