OCUSAN DRY EYES- polyethylene glycol 400, propylene glycol liquid
DLC Laboratories, Inc.

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.

Ocusan ® Dry Eyes

Drug Facts

Active ingredients	***Purposes***
Polyethylene Glycol 400 0.4%	Lubricant
Proplyene Glycol 0.3%	Lubricant

Use

- for the temporary relief of burning and irritation due to dryness of the eye

Warnings

For external use only

Do Not Use

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

When using this product

- do not touch tip of container to any surface to avoid contamination
- 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 of the eye
- condition worsens or lasts more than 72 hours

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

Directions

- shake well before using
- instill 1 to 2 drops in the affected eye(s) as needed

Other information

- store at room temperature

Inactive ingredients

benzalkonium chloride, boric acid, hydrochloric acid, hypromellose, potassium chloride, sodium chloride, sodium hydroxide, water for injection

Questions

1-800-858-3889

Distributed by:
DLC LABORATORIES, INC.
PARAMOUNT, CA 90723 USA

PRINCIPAL DISPLAY PANEL - 15 mL Bottle Box

Advanced Formula!

Fase Acting

Ocusan ®

Dry Eyes

Moisturizes & Soothes Dry Eyes

Lubricant Eye Drops / Relieves Buring and Irritation

STERILE

1/2 FL OZ (15 mL)

OCUSAN DRY EYES

polyethylene glycol 400, propylene glycol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:24286-1292
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	3 mg in 1 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ, POLYETHYLENE GLYCOL, UNSPECIFIED - UNII:3WQ0SDW1A)	POLYETHYLENE GLYCOL 400	4 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:24286-1292-5	1 in 1 BOX	12/16/2019	
1		15 mL in 1 BOTTLE; 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	12/16/2019	

Labeler - DLC Laboratories, Inc. (093351930)

Revised: 1/2023 DLC Laboratories, Inc.