

MINIDROPS- lubricant eye drops liquid
Optics Laboratory, 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.

ACTIVE INGREDIENTS SECTION

POVIDONE 0.6%

POLYVINYL ALCOHOL 1.4%

PURPOSE SECTION

EYE LUBRICANT

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.

INDICATIONS AND USE SECTION

USES FOR TEMPORARY RELIEF OF BURNING AND IRRITATION DUE TO DRYNESS OF THE EYE

FOR USE AS A PROTECTANT AGAINST FURTHER IRRITATION OR TO RELIEVE DRYNESS OF THE EYE

WARNINGS SECTION

WARNINGS DO NOT USE IF THE SINGLE-DOSE DROPPER IS NOT INTACT. TO AVOID CONTAMINATION, DO NOT TOUCH TIP OF DROPPER TO ANY SURFACE. ONCE OPENED, DISCARD.

IF YOU EXPERIENCE EYE PAIN, CHANGES IN VISION, CONTINUED REDNESS OR IRRITATION OF THE EYE, OR IF THE CONDITIONS WORSENS OR PERSISTS FOR MORE THAN 72 HOURS, DISCONTINUE USE AND CONSULT A DOCTOR. IF SOLUTION CHANGES COLOR OR BECOMES CLOUDY, DO NOT USE

OTHER INFORMATION STORE AT ROOM TEMPERATURE 15-30 DEGREES C (59-86 DEGREES F)

DOSAGE AND ADMINISTRATION SECTION

DIRECTIONS TWIST OFF TAB COMPLETELY TO OPEN. INSTILL 1 TO 2 DROPS IN THE AFFECTED EYE(S) AS NEEDED. DISCARD DROPPER AND DO NOT REUSE.

INACTIVE INGREDIENTS SECTION

SODIUM CHLORIDE, WATER (PURIFIED)

QUESTIONS SECTION

QUESTIONS OR COMMENTS? 800-968-6788

PACKAGE LABEL

MINIDROPS LUBRICANT EYE DROPS DROPPER

INDIVIDUAL PACKAGE TO AVOID CROSS-CONTAMINATION FROM REPEATED USE

COMPACT MINI-SIZE EASY TO CARRY OPTICS MANUFACTURED FOR OPTICS
LABORATORY INC.EL MONTE, CA 91731 USA MADE IN TAIWAN

STERILE PRESERVATIVE FREE USE ONLY IF SINGLE-USE DROPPER IS INTACT USE AS A
PROTECTANT AGAINST FURTHER IRRITATION OR TO RELIEVE DRYNESS OF THE EYE
PRESERVATIVE-FREE FORMULA SAFE FOR DAILY USE SINGLE-USE DROPPERS,
CONVENIENT TO USE AND CARRY

MINIDROPS AND OPTIC LABORATORY ARE TRADEMARKS OF OPTIC LABORATORY INC

Drug Facts

Active Ingredients (in each ml)
 Polyvinyl Alcohol 1mg
 Purposone Drug
 Eye Lubricant

Purpose
 Eye Lubricant

Indications
 ■ for the temporary relief of burning and irritation due to dryness of the eye
 ■ for use as a lubricant to prevent further irritation or to relieve dryness of the eye

Warnings
 Do not use if the single-use dropper is not intact. To avoid contamination, do not touch tip of dropper to any surface.
 Once opened, discard.
 If you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the conditions worsens or persists for more than 72 hours, discontinue use and consult a doctor. If solution changes color or becomes cloudy, do not use.
Keep out of reach of children. If swallowed, get medical help or contact a poison control center right away.

Manufactured for: Optics Laboratory, Inc. El Monte, CA 91731 USA Made in Taiwan
 MiniDrops and Optics Laboratory are trademarks of Optics Laboratory, Inc. © Optics Laboratory, Inc.
 www.minidrops.com

OPTICS
 LABORATORY

SAMPLE

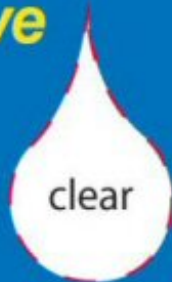
MiniDrops[®]

Eye Therapy

Lubricant Eye Drops

Preservative Free

*Soothing Relief
 For Eyes*



5 Sterile Single-Use Droppers

0.02 FL OZ EA
 (0.5) ml EA

- Relieves dry eyes, discomfort and minor irritation resulting from lengthy driving time and overexposure to heaters and air conditioning
- Preservative free formula recommended for safe daily use.
- Moisturizes your eyes with lasting lubrication like natural tears.

Directions:
 ■ Twist off tab completely to open
 ■ Instill 1 to 2 drops in the affected eye(s) as needed
 ■ Discard dropper and do not reuse

Other Information
 Store at room temperature
 15-30°C (59-86°F)

Inactive Ingredient
 Sodium Chloride

Questions or comments? 800-968-6788

EXP:
 LOT:

Drug Facts

Active Ingredients (in each ml) **Purpose**
 Povidone 0.6% Eye Lubricant
 Polyvinyl Alcohol 1.4% Eye Lubricant

Uses
 ■ for the temporary relief of burning and irritation due to dryness of the eye
 ■ for use as a protectant against further irritation or to relieve dryness of the eye

Warnings
 Do not use if the single-use dropper is not intact. To avoid contamination, do not touch tip of dropper to any surface. Once opened, discard.
 If you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the conditions worsens or persists for more than 72 hours, discontinue use and consult a doctor. If solution changes color or becomes cloudy, do not use.
 Keep out of reach of children. If swallowed, get medical help or contact a poison Control Center right away.

Directions:
 ■ Twist off tab completely to open
 ■ Instill 1 to 2 drops in the affected eye(s) as needed
 ■ Discard dropper and do not reuse

Other Information Store at room temperature
 15-30°C (59-86°F)

Inactive Ingredients Sodium Chloride,
 Water (Purified)

Questions or comments? 800-968-6788

MiniDrops®
Lubricant Eye Drops

OPTICS®
MiniDrops®
Lubricant Eye Drops

OPTICS®
MiniDrops®
Lubricant Eye Drops

Sterile Preservative Free

Sterile Preservative Free

30 Sterile Single-Use Droppers
0.02 FL OZ EA
(0.5) ml EA

Use only if single-use dropper is intact.

www.minidrops.com

res

MINIDROPS

lubricant eye drops liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64108-212
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE, UNSPECIFIED	6.0 mg in 1 mL
POLYVINYL ALCOHOL (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	POLYVINYL ALCOHOL	14.0 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	.9 mg in 1 mL
WATER (UNII: 059QF0K00R)	97.1 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64108-212-13	5 in 1 BOX	04/01/1991	
1	NDC:64108-212-10	0.5 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:64108-212-12	30 in 1 BOX	04/01/1991	
2	NDC:64108-212-10	0.5 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
	NDC:64108-212			

3	NDC:64108-212-65	65 in 1 BOX	04/01/1991	
3	NDC:64108-212-10	0.5 mL in 1 VIAL, SINGLE-USE; 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	04/01/1991	

Labeler - Optics Laboratory, Inc (018503552)

Registrant - Optics Laboratory, Inc (018503552)

Revised: 2/2019

Optics Laboratory, Inc