REFRESH LACRI-LUBE- mineral oil, petrolatum ointment
Allergan, Inc.

REFRESH® LACRI-LUBE®
Drug Facts

Active ingredients

Mineral Oil 42.5%

White Petrolatum 56.8%

Purpose

Eye lubricant

Eye lubricant

Uses

- For the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun.
- May be used as a protectant against further irritation.

Warnings

- **For external use only.**
- **To avoid contamination, do not touch tip of container to any surface.**
- **Replace cap after using.**

Stop use and ask a doctor if

you experience eye pain, changes in vision, continued redness or irritation of the eye, or if the 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

Pull down the lower lid of the affected eye and apply a small amount (one-fourth inch) of ointment to the inside of the eyelid.

Other information

- Store away from heat.
- Protect from freezing.
- Use only if tape seals on top and bottom flaps are intact.

- Use before expiration date marked on container.
- Store at 59°-77°F (15°-25°C).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Chlorobutanol and Lanolin alcohols.

Questions or comments?

1.800.678.1605

refreshbrand.com

Principal Display Panel

NDC 0023-0312-04

Lubricant Eye Ointment

Refresh®
LACRI-LUBE®

Nighttime Relief for
Intense Eye Dryness

Net wt. 0.12 oz (3.5 g) Sterile

REFRESH LACRI-LUBE

mineral oil, petrolatum ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0312
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	425 mg in 1 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	568 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CHLOROBUTANOL (UNII: HM4YQM8WRC)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0312-04	1 in 1 CARTON	12/15/1977	
1		3.5 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0023-0312-07	1 in 1 CARTON	12/15/1977	
2		7 g in 1 TUBE; 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	12/15/1977	

Labeler - Allergan, Inc. (144796497)

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