

REFRESH LACRI-LUBE- mineral oil, petrolatum ointment

Allergan, 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.

Drug Facts

Active ingredients

Mineral Oil 42.5%

White Petrolatum 56.8%

Purpose

Eye lubricant

Eye lubricant

Uses

- For use as a protectant against further irritation or to relieve dryness of the eye.

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

- Use only if imprinted tape seals on top and bottom flaps are intact and clearly legible.
- Store away from heat. Protect from freezing.
- Use before expiration date marked on container.
- Store at 59°-86°F (15°-30°C)
- **RETAIN THIS CARTON FOR FUTURE REFERENCE.**

Inactive ingredients

Chlorobutanol and lanolin alcohols.

Questions or comments?

☎ 1.800.678.1605

refreshbrand.com

HOW SUPPLIED: 0.12 oz (3.5 g) and 0.25 oz (7 g)

Principal Display Panel

Refresh[®]

Lacri-Lube[®]

Lubricant Eye Ointment

Ointment

for nighttime

dry eye relief

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 FINAL	part349	12/15/1977	

Labeler - Allergan, Inc. (144796497)

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Allergan, Inc.