REFRESH P.M.- 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.

REFRESH P.M.® (Preservative-free)
Drug Facts

Active ingredients

Mineral Oil 42.5%

White Petrolatum 57.3%

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

Lanolin alcohols.

Questions or comments?

1.800.678.1605

refreshbrand.com

v1.0DFL0667

PRINCIPAL DISPLAY PANEL

NDC 0023-0667-04
PRESERVE-FREE
Lubricant Eye Ointment
For Eye Dryness
Refresh
P.M.®
Nighttime Relief for
Intense Eye Dryness
Net wt. 0.12 oz (3.5 g) Sterile

PRINCIPAL DISPLAY PANEL

NDC 0023-0240-04
Refresh
P.M.
Lubricant Eye Ointment
NIGHTTIME OINTMENT
Net wt. 0.12 oz (3.5 g) Sterile

Image Not Available

REFRESH P.M.

mineral oil, petrolatum ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0240
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	573 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
lanolin alcohols (UNII: 884C3FA9HE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0240-21	1 in 1 CARTON	04/23/1990	04/17/2019
1		3.5 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0023-0240-04	1 in 1 CARTON	04/23/1990	04/17/2019
2		3.5 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	04/23/1990	

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Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0667
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
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Inactive Ingredients

Ingredient Name	Strength
lanolin alcohols (UNII: 884C3FA9HE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0667-04	1 in 1 CARTON	08/26/2016	
1		3.5 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	08/26/2016	

Labeler - Allergan, Inc. (144796497)

Revised: 6/2022 Allergan, Inc.