LUBRICANT PM- mineral oil, white petrolatum lubricant eye ointment ointment AACE PHARMACEUTICALS, INC.

SPL UNCLASSIFIED SECTION

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

Mineral Oil 42.5% and White Petrolatum 57.3%

PURPOSE

Lubricant

USES

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

WARNINGS

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

Stop use and ask a doctor if

- you feel eye pain.
- changes in vision occur.
- redness or irritation of the eye(s) gets worse, persists 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

• 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 15° to 30°C (59° to 86°F).

RETAIN THIS CARTON FOR FUTURE REFERENCE.

INACTIVE INGREDIENTS

Lanolin Alcohols.

PRINCIPAL DISPLAY PANEL

NDC#: 71406-124-35

Compare to Refresh P.M. ® Ointment active ingredients*

Lubricant PM Ointment

Sensitive

Preservative Free

Ointment for Nighttime Dry Eye Relief

Net Wt. 3.5 g (1/8 oz.) Sterile

Distributed By:

AACE Pharmaceuticals, Inc., Fairfield, NJ 07004

www.aacepharma.com

Rev. 02

*This product is not manufactured by Allergan, Inc. distributor of Refresh PM $^{\circledR}$ Ointment.



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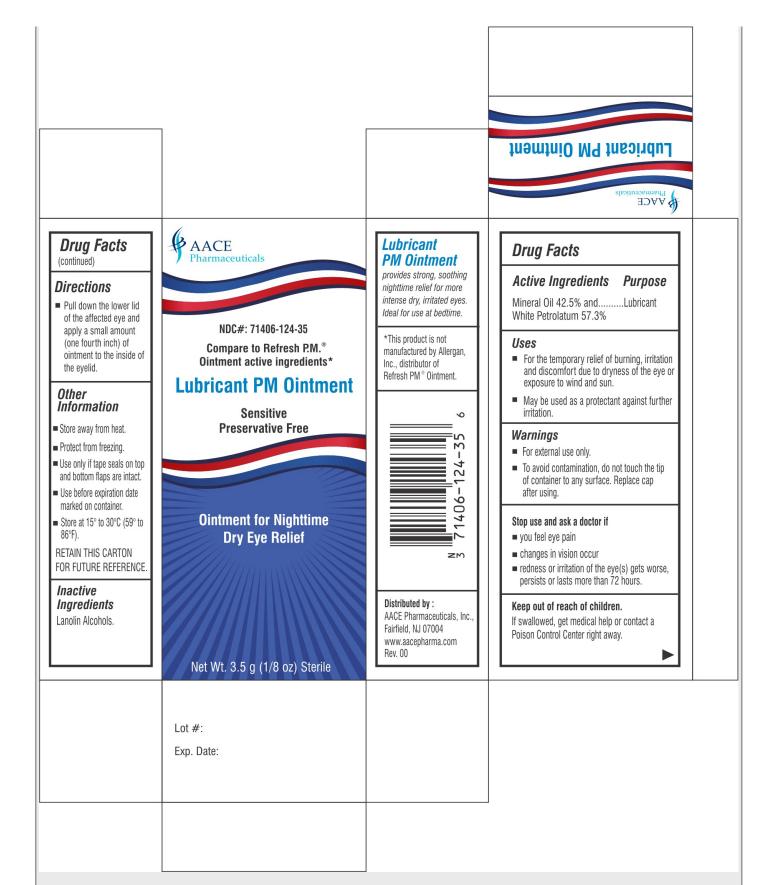
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71406-124		
Route of Administration	OPHTHALMIC				

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)				

Product Characteristics				
Color	Score			
Shape	Size			
Flavor	Imprint Code			
Contains				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:71406-124- 35	1 in 1 CARTON	03/19/2020		
1		3.5 g in 1 TUBE; Type 0: Not a Combination Product			



Marketing InformationMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateOTC Monograph DrugM01803/19/2020

Labeler - AACE PHARMACEUTICALS, INC. (080630748)

Revised: 1/2024 AACE PHARMACEUTICALS, INC.