

LUBRICANT EYE- mineral oil and petrolatum ointment

Akorn

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 ingredient

Mineral Oil

12%

White Petrolatum

88%

Contains no preservatives or lanolin.

Purpose

Eye lubricant

Eye lubricant

Uses

For use as a lubricant for the temporary relief of burning and irritation due to dryness of the eyes.

Warnings

- **Do not use if bottom ridge of tube is exposed.**
- **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 the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Pull down lower lid of the affected eye(s) and apply small amount (one-fourth inch) of ointment to the inside of the eyelid, one or more times daily, or as directed by a doctor.

Other information

- **Store at 20° to 25°C (68° to 77°F)**
[see USP Controlled Room Temperature].
- **Store away from heat.**
- **Protect from freezing.**
- **Keep tightly closed.**
- **See crimp for Control Number and Expiration Date.**
- **RETAIN THIS CARTON FOR FUTURE REFERENCE.**

NJTROAC Rev. 06/12

Principal Display Panel Text for Container Label:

NDC 17478-063-35 Akorn Logo

Lubricant Eye OINTMENT

Nighttime Relief For Dry Eyes Sterile

FOR OPHTHALMIC USE ONLY Net Wt. 3.5 g (1/8oz.)

NDC 17478-063-35 

Lubricant Eye OINTMENT

Nighttime Relief for Dry Eyes **Sterile**

FOR OPHTHALMIC USE ONLY **Net Wt. 3.5 g (1/8 oz.)**

READ OUTER CARTON FOR INFORMATION BEFORE USING.

Active ingredient: Mineral Oil 12%, White Petrolatum 88%.
Contains no preservatives or lanolin.

Directions: Pull down the lower lid of the affected eye(s) and apply a small amount (one-fourth inch) of ointment to the inside of the eyelid, one or more times daily, or as directed by a doctor. **Uses:** For use as a lubricant for the temporary relief of burning and irritation due to dryness of the eyes. See crimp of tube for Control Number and Expiration Date.

Storage: Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature]. Keep tightly closed.

WARNING: KEEP OUT OF THE REACH OF CHILDREN.

DO NOT USE IF BOTTOM RIDGE OF TUBE CAP IS EXPOSED.

Manufactured by:
Akorn, Inc., Lake Forest, IL 60045
www.akorn.com NJTROAL Rev. 06/12 (01) 00317478063353

LT-0819-5 



Principal Display Panel Text for Carton Label:

NDC 17478-063-35

3.5 g

Lubricant

Eye

OINTMENT

Nighttime Relief

For Dry Eyes

Nighttime Relief

Plus Protection

Sterile

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



0705

Drug Facts

Active ingredient Purpose

Mineral Oil
12%.....Eye lubricant
White Petrolatum
88%.....Eye lubricant
Contains no preservatives or lanolin.

Uses For use as a lubricant for the temporary relief of burning and irritation due to dryness of the eyes.

Warnings

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- Replace cap after using.

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Drug Facts

(continued)

Directions

Pull down lower lid of the affected eye(s) and apply small amount (one-fourth inch) of ointment to the inside of the eyelid, one or more times daily, or as directed by a doctor.

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NDC 17478-063-35

FOR OPHTHALMIC USE ONLY

3.5 g

Lubricant Eye OINTMENT

Nighttime Relief for Dry Eyes



Nighttime Relief Plus Protection

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

Questions
1-800-932-5676



3 17478-063-35 3

AKORN

Manufactured by:
Akorn, Inc.
Lake Forest, IL 60045
www.akorn.com

NJTROAC Rev. 06/12

mineral oil and petrolatum ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17478-063
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Mineral Oil (UNII: T5L8T28FGP) (Mineral Oil - UNII:T5L8T28FGP)	Mineral Oil	120 mg in 1 g
Petrolatum (UNII: 4T6H12BN9U) (Petrolatum - UNII:4T6H12BN9U)	Petrolatum	880 mg in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17478-063-35	1 in 1 CARTON	04/01/1993	
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	04/01/1993	

Labeler - Akorn (117696770)

Registrant - Akorn Operating Company LLC (117693100)

Establishment

Name	Address	ID/FEI	Business Operations
Akorn		117696840	MANUFACTURE(17478-063) , ANALYSIS(17478-063) , STERILIZE(17478-063) , PACK(17478-063) , LABEL(17478-063)

Revised: 1/2022

Akorn