

DRY EYES- white petrolatum mineral oil ointment
Cardinal Health

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.

Artificial Tears Ointment Drug Facts

Active ingredients

Mineral oil 15%
White petrolatum 83%

Purpose

Lubricant

Uses

- to prevent further irritation
- to relieve dryness of the eye

Warnings

When using this product

- replace cap after use
- do not touch tip of container to any surface to avoid contamination

Stop use and ask a doctor if

- condition worsens or persists for more than 72 hours
- you experience eye pain, changes in vision, continued redness or irritation of the eye

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(s)
apply a small amount (1/4 inch) of ointment to the inside of eyelid
apply one or more times daily or as directed by a doctor

Other information

- store at 15° - 30°C (59° - 86°F)
- keep tightly closed
- see crimp of tube or carton for Lot Number and Expiration Date

Inactive ingredient

lanolin oil

Questions?

Serious side effects associated with use of this product may be reported to

1-800-323-0000

DO NOT USE IF CAP AND NECKRING ARE NOT INTACT.

Package/Label Principal Display Panel



NDC 37205-134-79

LEADER®

Compare to Refresh P.M. active ingredient*

Dry Eyes

Lubricant Eye Ointment
(Sterile)

Preservative Free

Prevents irritation & relieves dryness of the eye

SATISFACTION GUARANTEED Net wt. 1/8 oz. (3.5 g)

DRY EYES

white petrolatum mineral oil ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37205-134
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	150 mg in 1 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	830 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
LANOLIN (UNII: 7EV65EAW6H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37205-134-79	3.5 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	11/15/2011	

Labeler - Cardinal Health (097537435)

Registrant - Bausch & Lomb Incorporated (196603781)

Establishment

Name	Address	ID/FEI	Business Operations
Bausch & Lomb Incorporated		807927397	MANUFACTURE

