NIGHTTIME DRY-EYE RELIEF- mineral oil, and white petrolatum ointment CVS

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.

Nighttime Dry-Eye Relief

Active ingredients Purpose

Mineral oil 42.5%..... Eye Lubricant

White petrolatum 57.3%..... Eye Lubricant

Uses

- For the temporary relief of burning, irritation, and discomfort due to dryness or 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.
- Keep tube tightly closed.

Stop use and ask a doctor if:

- You experience eye pain, changes in vision, continued redness or irritation of the eye.
- 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 at 1-800-222-1222.

Directions

- Wash your hands.
- Pull down the lower lid of the affected eye and gently apply a small amount (1/4) of ointment to the inside of the eyelid.

Other information

- store away from heat
- protect from freezing
- use before expiration date on the tube
- store at 59°-86° (15°-30°)

Inactive ingredient

lanolin alcohol

Directions

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Nighttime Dry-Eye Relief Lubricant eye ontment

Preservative Free

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NDC # 59779-568-13

Preservative Free

Nighttime Dry-Eye Relief

LUBRICANT EYE OINTMENT

- Sensitive formula
- Protects & moisturizes at bedtime
- Soothes & prevents irritation for dry, sensitive eyes

Actual Product Size on Side Panel

STERILE NET WT 0.12 OZ (3.5 g) DO NOT USE IF CAP NECK SEAL IS CUT, BROKEN OR MISSING.

Save carton for complete drug facts.

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CVS Quality

V-19655

Package Contains One Tube

Actual Size

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Drug Facts (continued)

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NIGHTTIME DRY-EYE RELIEF

mineral oil, and white petrolatum ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-568
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:69842-568-13	1 in 1 BOX	08/30/2019		
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	08/30/2019		
		o o	Will be this	

Labeler - CVS (062312574)

Revised: 8/2019 CVS