NIGHTIME RELIEF LUBRICANT EYE- mineral oil, and white petrolatum ointment
Target Corporation

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 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 contaainer to any surface. Replace cap after using.

Stop use and ask a doctor if you experience eye pain, changes in vision, continued reness or irritation of the eye, or if the condition worsens or persists 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 (1/4") of ointment to the inside of the eyelid

Other information

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

Inactive ingredient

lanolin alcohol

DISTRIBUTED BY:

TARGET CORP.

MINNEAPOLIS, MN 55403

MADE IN KOREA

NIGHTIME RELIEF LUBRICANT EYE

mineral oil, and white petrolatum ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-575
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	533 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:11673-575-12	1 in 1 BOX		
1		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	03/05/2015	

Labeler - Target Corporation (006961700)

Revised: 3/2015 Target Corporation