

ALTALUBE- mineral oil and petrolatum ointment
Altaire Pharmaceuticals Inc.

Altalube Ointment

ActivEyes

Altalube Ointment 3.5g

NDC 59390-198-50

Drug Facts

Active ingredients

Mineral Oil 15% and White Petrolatum 85%

Purpose

Eye Lubricant

Uses

- as a lubricant to prevent further irritation
- to relieve dryness of the eye(s).

Warnings

- Save box for complete information.
- **For use in the eyes only.**

When using this product

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

Stop use and ask a doctor if

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

Keep this and all drugs 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 and apply a small amount (one-fourth inch) of ointment to the inside of the eyelid every 3-4 hours or as directed by a doctor.

Other information

- store at room temperature 15°-30°C (59°-86°F). • protect from freezing.
- see crimp of tube or box for lot number and expiration date.
- keep tightly closed.
- does not contain preservatives.

Inactive Ingredients

None remove

Questions or comments?

- (631) 722-5988 • 9AM - 5 PM EST Monday- Friday

PRINCIPAL DISPLAY PANEL

ActiEyes Altalube Ointment White Petrolatum and Mineral Oil Lubricant Eye Ointment
Preservative Free Sterile NET WT 3.5g (1/8 OZ)



ALTALUBE

mineral oil and petrolatum ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59390-198
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	850 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:59390-198-50	1 in 1 CARTON	02/01/2002	
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 Drug	M018	02/01/2002	

Labeler - Altaire Pharmaceuticals Inc. (786790378)**Registrant** - Altaire Pharmaceuticals, Inc. (786790378)**Establishment**

Name	Address	ID/FEI	Business Operations
Altaire Pharmaceuticals, Inc.		786790378	manufacture(59390-198)

Revised: 12/2023

Altaire Pharmaceuticals Inc.