REFRESH CLASSIC- polyvinyl alcohol, povidone solution/ drops
Allergan, Inc.

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.

REFRESH Classic

Drug Facts

Active ingredients

Polyvinyl Alcohol 1.4%

Povidone 0.6%

Purpose

Eye lubricant

Eye lubricant

Uses

- For the temporary relief of burning, irritation, and discomfort due to dryness of 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. Do not reuse. Once opened, discard.**
- **Do not touch unit-dose tip to eye.**
- **If solution changes color or becomes cloudy, do not use.**

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 reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

To open, **TWIST AND PULL TAB TO REMOVE**. Instill 1 or 2 drops in the affected eye(s) as needed and discard container.

Other information

- Use only if single-use container is intact.
- Use before expiration date marked on container.
- Store at 59°-86°F (15°-30°C).
- RETAIN THIS CARTON FOR FUTURE REFERENCE.

Inactive ingredients

Purified water and sodium chloride. May also contain hydrochloric acid and/or sodium hydroxide to adjust pH.

Questions or comments?

1.800.433.8871

refreshbrand.com

NDC 0023-0506-01

Preservative-free

***Refresh*®**
Classic
Lubricant Eye Drops

Moisture drops
for dry eyes

30 Single-Use Containers
0.01 fl oz (0.4 mL) each Sterile

REFRESH CLASSIC

polyvinyl alcohol, povidone solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0506
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Polyvinyl Alcohol (UNII: 532B59J990) (Polyvinyl Alcohol - UNII:532B59J990)	Polyvinyl Alcohol	14 mg in 1 mL
Povidone (UNII: FZ989GH94E) (Povidone - UNII:FZ989GH94E)	Povidone	6 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
sodium chloride (UNII: 451W47IQ8X)	
hydrochloric acid (UNII: QTT17582CB)	
sodium hydroxide (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0506-01	30 in 1 CARTON	09/12/1985	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:0023-0506-50	50 in 1 CARTON	09/12/1985	
2		0.4 mL in 1 VIAL, SINGLE-USE; 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	09/12/1985	

Labeler - Allergan, Inc. (144796497)

Establishment

Name	Address	ID/FEI	Business Operations
Allergan Sales, LLC		362898611	MANUFACTURE(0023-0506)

Establishment

Name	Address	ID/FEI	Business Operations
Allergan Pharmaceuticals Ireland		219682291	MANUFACTURE(0023-0506)

Revised: 4/2011 Allergan, Inc.