BLINK GEL TEARS- polyethylene glycol 400 solution/ drops
Johnson & Johnson Surgical Vision, 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.

Drug Facts

OTC - ACTIVE INGREDIENT SECTION

Polyethylene Glycol 400 0.25%

Purpose

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. Replace cap after using.
- Do not use if solution changes color or becomes cloudy.

Stop use and ask 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.

OTC - KEEP OUT OF REACH OF CHILDREN SECTION

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

Directions

Instill 1 or 2 drops in the affected eye(s) as needed.

Other Information

Use only if tape seals on top and bottom flaps are intact.

RETAIN THIS CARTON FOR FUTURE REFERENCE.

INACTIVE INGREDIENT SECTION

Boric acid, calcium chloride, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, sodium chlorite (OcuPure® Brand) as a preservative; Sodium Hyaluronate.

OTC - QUESTIONS SECTION

In the U.S. call 1-800-347-5005

www.yourhealthyeyes.com

PRINCIPAL DISPLAY PANEL - 10 mL Bottle Carton

For both Daytime & Nighttime Relief

Relief with every blink

blink
gel Tears®

Lubricating Eye Drops

Moderate-Severe Dry Eye

0.34 FL OZ
(10 mL) STERILE

PRINCIPAL DISPLAY PANEL - 2 mL Bottle Carton

Relief with every blink

blink
gel Tears®

Lubricating Eye Drops

Moderate-Severe Dry Eye

PROFESSIONAL USE – NOT FOR RESALE

0.06 FL OZ

(2 mL) STERILE

BLINK GEL TEARS

polyethylene glycol 400 solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:29943-004
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Polyethylene Glycol 400 (UNII: B697894SGQ) (Polyethylene Glycol 400 - UNII:B697894SGQ)	Polyethylene Glycol 400	2.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Boric Acid (UNII: R57ZHV85D4)	
Calcium Chloride (UNII: M4I0D6VV5M)	
Magnesium Chloride (UNII: 02F3473H9O)	
Potassium Chloride (UNII: 660YQ98I10)	
Water (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
Sodium Chloride (UNII: 451W47IQ8X)	
SODIUM CHLORITE (UNII: G538EBV4VF)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29943-004-10	1 in 1 CARTON	05/01/2008	
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:29943-004-02	1 in 1 CARTON	05/01/2008	
2		2 mL in 1 BOTTLE, DROPPER; 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	05/01/2008	

Labeler - Johnson & Johnson Surgical Vision, Inc. (103021940)