

**TEARS LUBRICANT- glycerin, hypromelloses, polyethylene glycol 400 liquid
Kareway Product, 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.

Pure-Aid Artificial Tears

Active Ingredient

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene Glycol 400 1%

Purpose

Lubricant

Lubricant

Lubricant

Use

- for the temporary relief of burning and irritation due to dryness of the eye
- for protection against further irritation

Warnings

When using this product

- remove contact lenses before using
- do not use if this solution changes color or become cloudy
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye lasts
- condition worsens or lasts more than 72 hours

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

- Put 1 to 2 drops in the affected eye(s) up to 4 times daily

- children under 6 years of age: ask a doctor

Other information

- store between 15° to 25°C (59°F to 77°F)

Inactive ingredients

Ascorbic acid, benzalkonium chloride, boric acid, dextrose, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, sodium citrate, sodium lactate

Package principal display panel

Pureaid Artificial Tears

Moisturizes and Soothes dry eyes
• Protects against further irritation

Caution: Do not use if the tamper-resistant bottle cap is broken or opened.

Pure-Aid™
Eye Drops STERILE
TEARS

NDC No. 47510-0043-5

Compare to Visine®
Tears Eye Drops™

Eye Drops STERILE

TEARS
LUBRICANT EYE DROPS

Dry eye relief
formula

1/2 FL OZ (15mL)

Pure-Aid™

Eye Drops STERILE

TEARS
LUBRICANT EYE DROPS

KAREWAY

Exclusively distributed by:
Kareway Product Inc.
2550 S. Dominguez Hills Dr.
Compton, CA 90220
Made in Korea

*This product is not manufactured or distributed by Johnson & Johnson Healthcare Products Division of McNEIL-PPC, Inc.

Drug Facts

Active ingredients	Purpose
Glycerin 0.2%	Lubricant
Hypromellose 0.2%	Lubricant
Polyethylene Glycol 400 1%	Lubricant

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If pregnant or breast-feeding, ask a health professional before use.

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

Drug Facts (continued)

Directions

- put 1 to 2 drops in the affected eye(s) as needed
- children under 6 years of age: ask a doctor

Other information

- store at 15° to 30°C (59° to 86°F)

Inactive ingredients

ascorbic acid, benzalkonium chloride, boric acid, glucose, glycine, monobasic sodium phosphate monohydrate, potassium chloride, purified water, sodium borate, sodium chloride, sodium citrate hydrate, sodium hydroxide



LOT:

EXP.

TEARS LUBRICANT

glycerin, hypromelloses, polyethylene glycol 400 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67510-0063
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2 mg in 1 mL
HYPROMELLOSES (UNII: 3NXW29V3WO) (HYPROMELLOSES - UNII:3NXW29V3WO)	HYPROMELLOSES	2 mg in 1 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67510-0063-5	1 in 1 BOX	05/30/2018	
1		15 mL in 1 BOTTLE; 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	05/30/2018	

Labeler - Kareway Product, Inc. (121840057)