

VISINE DRY EYE RELIEF- polyethylene glycol 400 solution/ drops
Kenvue Brands LLC

VISINE® Dry Eye Relief All Day Comfort

Drug Facts

Active ingredient

Polyethylene glycol 400 1%

Purpose

Lubricant

Uses

- for the temporary relief of burning, irritation, and discomfort due to dryness of the eye or exposure to wind or sun
- for protection against further irritation

Warnings

For external use only

When using this product

- remove contact lenses before using
- do not use if this solution changes color or becomes 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

- adults and children 6 years of age and over: put 1 or 2 drops in the affected eye(s) as needed
- children under 6 years of age: consult a doctor

Other information

Store at room temperature

Inactive ingredients

ascorbic acid, boric acid, dextrose, glycerin, glycine, hypromellose, magnesium chloride, polyquaternium-42, potassium chloride, purified water, sodium borate, sodium chloride, sodium citrate, sodium lactate, sodium phosphate dibasic

Questions?

call toll-free **888-734-7648** or **215-273-8755** (collect)

Distributed by: **JOHNSON & JOHNSON CONSUMER INC.** Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton

***UPTO 10HRS COMFORT**

VISINE®

DRY EYE RELIEF

ALL DAY*

COMFORT

LUBRICANT EYE DROPS

STARTS TO RELIEVE

DRY EYE IN SECONDS

Protects Against

Further Irritation

STERILE

1/2 FL OZ (15 mL)



VISINE DRY EYE RELIEF

polyethylene glycol 400 solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0365
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 -	POLYETHYLENE	10 mg

UNII:B697894SGQ, POLYETHYLENE GLYCOL, UNSPECIFIED - UNII:3WJQ0SDW1A) GLYCOL 400 in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
BORIC ACID (UNII: R57ZHV85D4)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCINE (UNII: TE7660XO1C)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POLIXETONIUM CHLORIDE (UNII: RNW5AM67C0)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0365-1	1 in 1 CARTON	04/27/2020	
1		15 mL in 1 BOTTLE, DROPPER; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	04/27/2020	

Labeler - Kenvue Brands LLC (118772437)

Revised: 11/2024

Kenvue Brands LLC