

**ARTIFICIAL TEARS LUBRICANT EYE DROPS- glycerin, hypromellose, polyethylene glycol 400 solution/ drops
KC Pharmaceuticals, Inc**

KC Pharmaceuticals, Inc (MFG) for Crocodile Tears (PLD)

Active ingredients

Glycerin 0.2%

Hypromellose 0.2%

Polyethylene glycol 1%

Purpose

Glycerin.....Lubricant

Hypromellose.....Lubricant

Polyethylene glycol 400.....Lubricant

Uses

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

Warnings

For external use only

Do not use this product if solution changes color or becomes cloudy

- to avoid contamination, do not touch tip of container to any
- remove contact lenses before using

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

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 (1-800-222-1222) right away.

Directions

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

Other information

Store at 15 °-30 °C (59 °-86 °F)

Inactive ingredients

benzalkonium chloride, dextrose, edetate disodium, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate, sodium phosphate dibasic, sodium

phosphate monobasic

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Artificial Tears
Sterile
CROCODILE TEARS™

Product of USA

Lubricant Eye Drops

Drug Facts

Active ingredients	Purposes
Glycerin 0.2%.....	Lubricant
Hypromellose 0.2%.....	Lubricant
Polyethylene glycol 400 1%.....	Lubricant

Uses

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

Warnings

For external use only

Do not use this product if solution changes color or becomes cloudy

When using this product

- to avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- remove contact lenses before using

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

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Directions

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Drug Facts (continued)

Other information
store at 15°-30°C (59°-86°F)


Inactive ingredients
benzalkonium chloride, dextrose, edetate disodium, potassium chloride, purified water, sodium bicarbonate, sodium chloride, sodium citrate, sodium phosphate dibasic, sodium phosphate monobasic

*This product is not manufactured or distributed by Johnson & Johnson Healthcare Products, distributor of Visine® Dry Eye Relief.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED NECKBAND IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

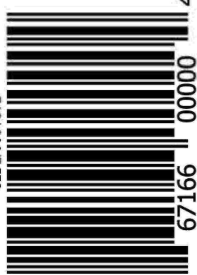
CROCODILE TEARS™



Product of USA

Artificial Tears
Lubricant Eye Drops

They don't sting.
They don't bite.



CED110651CTZ
8 67166 00000 4

Soothes Protects Moisturizes

0.5 fl oz (15mL) Sterile

LOT

EXP

Distributed by:
Crocodile Tears
Palm Beach Gardens, FL 33410
1-855-880-9538
www.croctears.com

ARTIFICIAL TEARS LUBRICANT EYE DROPS

glycerin, hypromellose, polyethylene glycol 400 solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55651-115
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.2 g in 100 mL
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) (HYPROMELLOSE, UNSPECIFIED - UNII:3NXW29V3WO)	HYPROMELLOSE, UNSPECIFIED	0.2 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE (UNII: IY9XDZ35W2)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-115-01	1 in 1 CARTON	03/25/2026	
1		15 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 Drug	M018	03/25/2026	

Labeler - KC Pharmaceuticals, Inc (174450460)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(55651-115) , pack(55651-115) , label(55651-115)

Revised: 3/2026

KC Pharmaceuticals, Inc