

**GOOD NEIGHBOR PHARMACY ARTIFICIAL TEARS LUBRICANT EYE DROPS-
polyvinyl alcohol, povidone solution/ drops
AmerisourceBergen**

GNP Artificial Tears (PLD)

Active ingredients

Polyvinyl alcohol.....0.5%

Povidone.....0.6%

Purpose

Eye lubricant

Uses

- for use as a protectant against further irritation or to relieve dryness of the eye
- for the temporary relief of discomfort due to minor irritations of the eye, or to exposure to wind or sun

Warnings

For external use only

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

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

When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using. Keep container tightly closed.
- remove contact lens before using

Keep out of the reach of children. If accidentally swallowed , get medical help or contact a Poison Control Center immediately.

Directions

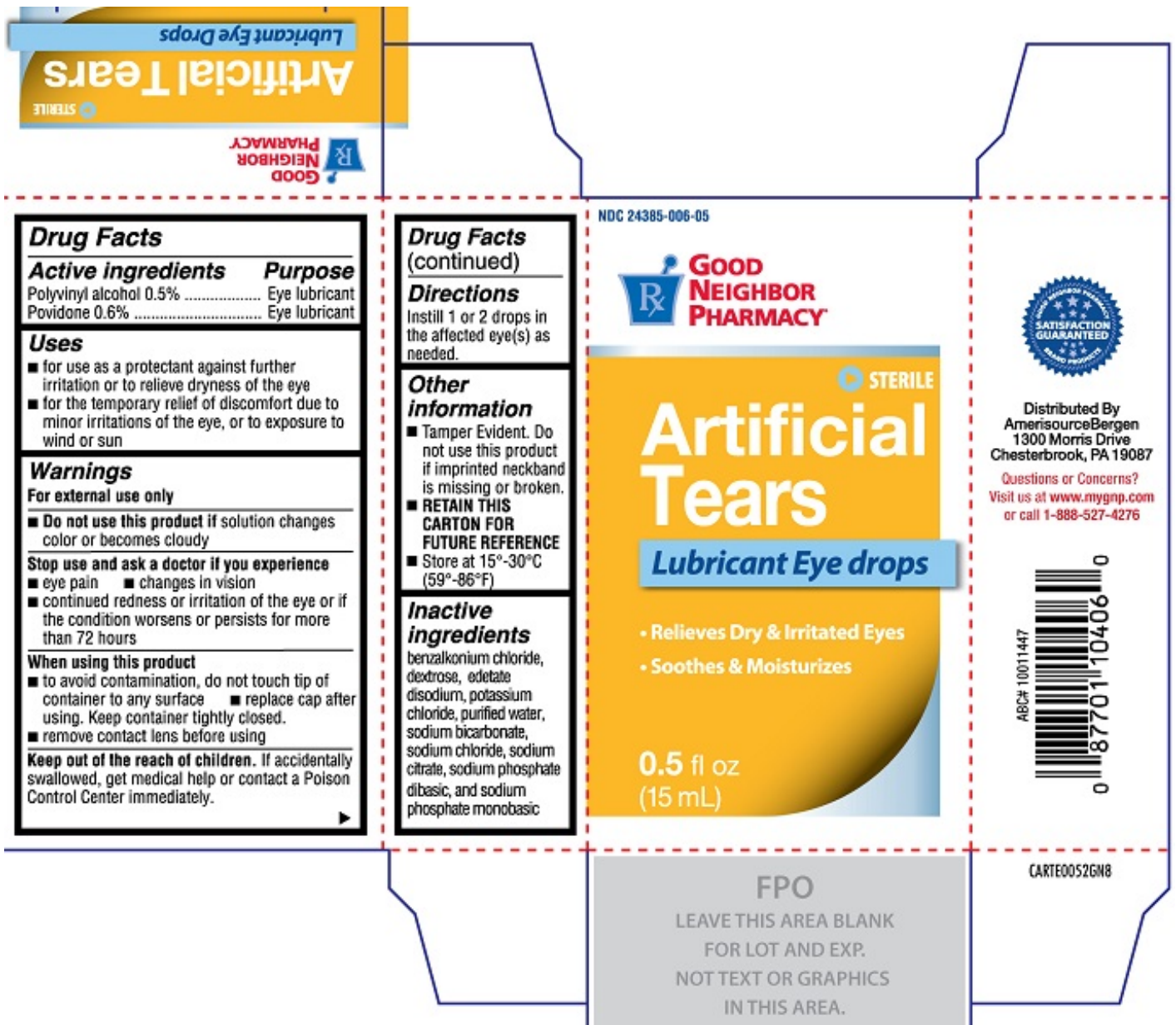
Instill 1 or 2 drops in the affected eye(s) as needed.

Other information

- Tamper Evident. Do not use this product if imprinted neckband is missing or broken.
- RETAIN THIS CARTON FOR FUTURE REFERENCE
- 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, and sodium phosphate monobasic



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24385-006
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL, UNSPECIFIED - UNII:532B59J990)	POLYVINYL ALCOHOL, UNSPECIFIED	0.5 g in 100 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	0.6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DEXTROSE (UNII: IY9XDZ35W2)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
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:24385-006-05	1 in 1 CARTON	01/29/2001	
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	01/29/2001	

Labeler - AmerisourceBergen (007914906)

Registrant - K.C. Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(24385-006) , pack(24385-006) , label(24385-006)

Revised: 12/2023

AmerisourceBergen