GOOD SENSE ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution
Geiss, Destin & Dunn, 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.

Good Sense Artificial Tears PLD

Active ingredients

Polyvinyl alcohol.....0.5%

Povidone.....0.6%

Purposes

Polyvinyl alcohol ...Eye lubricant

Povidone ...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**

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

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**

Directions

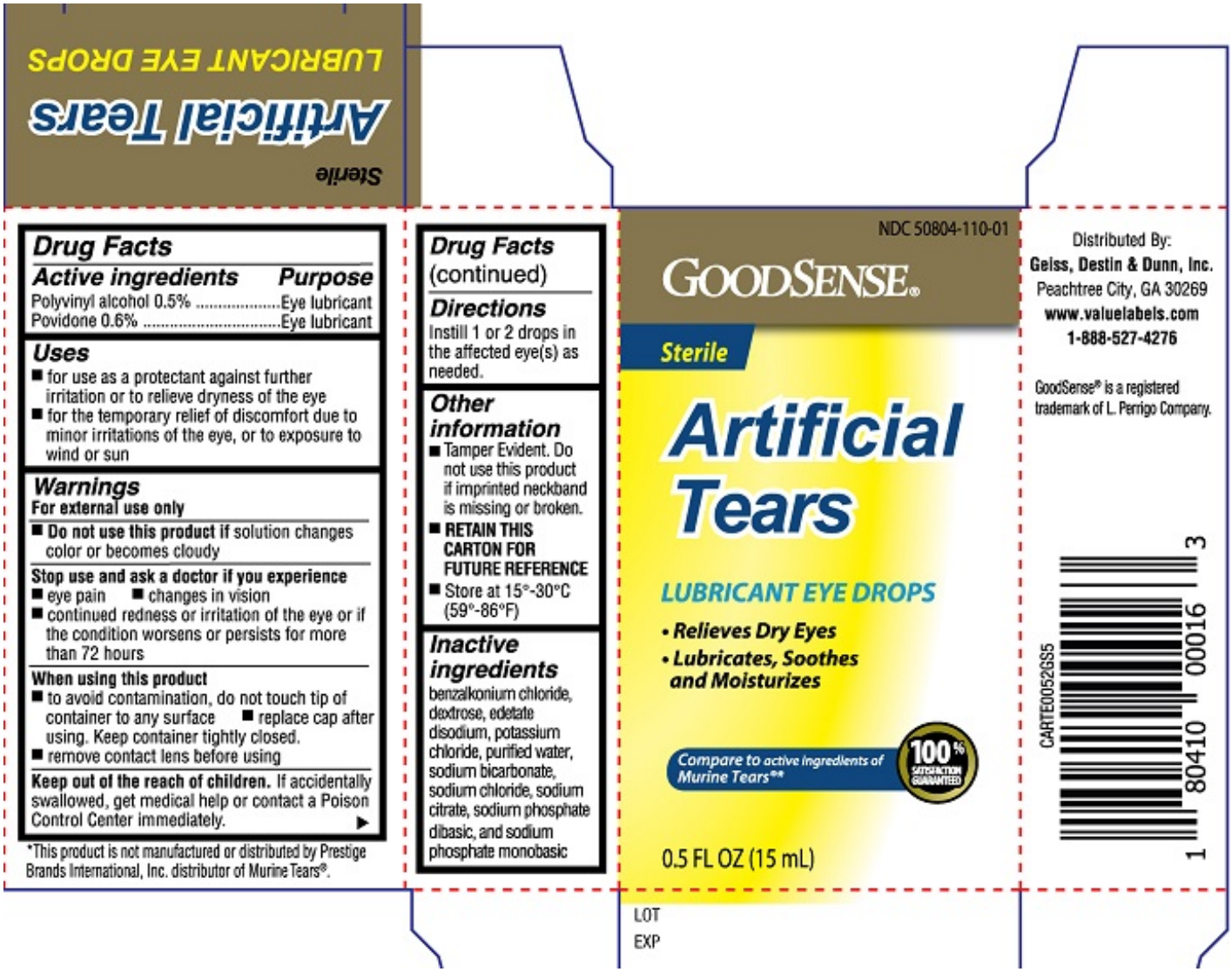
Instill 1 or 2 drops in the affected eye(s) as needed.

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



GOOD SENSE ARTIFICIAL TEARS

polyvinyl alcohol, povidone solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-110
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**
EDETATE DISODIUM (UNII: 7FLD91C86K)	

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-110-01	1 in 1 BOX	01/02/2020	
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 final	part349	01/02/2020	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(50804-110) , pack(50804-110) , label(50804-110)

Revised: 1/2020 Geiss, Destin & Dunn, Inc.