ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution NuCare Pharmaceuticals,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

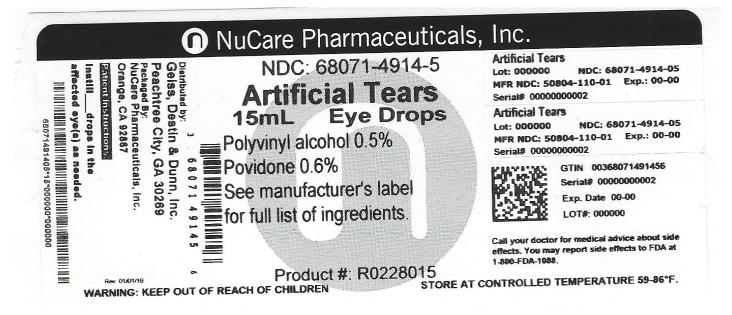
Instill 1 or 2 drops in the affected eye(s) as needed.

Other information

- Tamper Evident. Do not use this product if imprinted neckband is missing or broken.
- RETAIN THIS CARTON FOR FUTURE REFERENCE
- Store at 15º-30º C (59º-86ºF)

Inactive ingredients

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



ARTIFICIAL TEARS

polyvinyl alcohol, povidone solution

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4914(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: IY9XDZ 35W2)			

sc	SODIUM CITRATE (UNII: 1Q73Q2JULR)					
Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date		
1		15 mL in 1 BOX; Type 0: Not a Combination Product	01/14/2020			
Marketing Information						
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
ОТ	۲C monograph fina	l part349	01/02/2020			

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment				
Name	Address	ID/FEI	Business Operations	
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-4914)	

Revised: 2/2021

NuCare Pharmaceuticals, Inc.