QUALITY CHOICE ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution/drops

Chain Drug Marketing Association, Inc.

Quality Choice Artificial Tears 15 mL (PLD)

Active ingredients

Polyvinyl alcohol 0.5%

Povidone 0.6%

Purpose

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 productif 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



QUALITY CHOICE ARTIFICIAL TEARS

polyvinyl alcohol, povidone solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-223
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Basis of Strength	Ingredient Name	
DLYVINYL ALCOHOL 500 mg in 100 mL	POLYVINYL ALCOHOL (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	
OVIDONE 600 mg in 100 mL	POVIDONE (UNII: FZ 989GH94E) (POVIDONE - UNII:FZ 989GH94E)	
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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:63868-223-15	1 in 1 BOX	07/26/2019	
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	07/26/2019	

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(63868-223) , label(63868-223) , pack(63868-223)

Revised: 12/2023 Chain Drug Marketing Association, Inc.