

BEST CHOICE ARTIFICIAL TEARS- polyvinyl alcohol, povidone solution/ drops
Best Choice

Best 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 product** if solution changes color or becomes cloudy

When using this product

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

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 (1-800-222-1222) immediately.

Directions

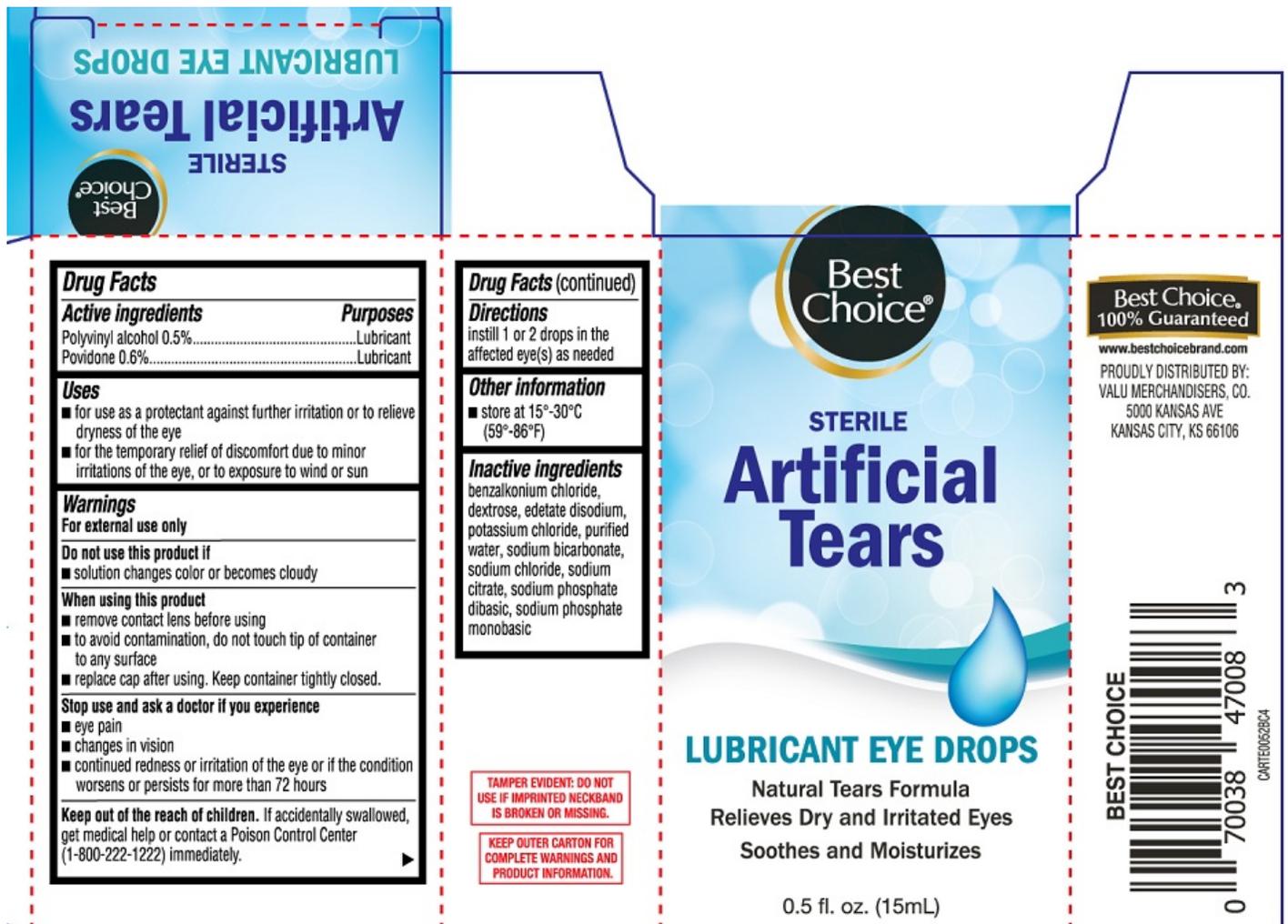
Instill 1 or 2 drops in the affected eye(s) as needed.

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, and sodium phosphate monobasic



BEST CHOICE ARTIFICIAL TEARS

polyvinyl alcohol, povidone solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-015
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYVINYL ALCOHOL (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	POLYVINYL ALCOHOL	0.5 g in 100 mL
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE	0.6 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:63941-015-01	1 in 1 BOX	03/21/2023	
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	03/21/2023	

Labeler - Best Choice (868703513)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(63941-015) , label(63941-015) , pack(63941-015)

Revised: 12/2025

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