

CLEAR VISION EYE DROPS- polyvinyl alcohol, povidone solution/ drops
Navajo Manufacturing Company 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.

Clear Vision Eye Drops

Drug Facts

Active ingredients

Polyvinyl alcohol 0.5%

Povidone 0.6%

Purpose

Lubricant

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).
- Keep card for complete warnings and product information.

Inactive ingredients

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

Questions or comments?

Call toll-free 1-800-525-5097

Package Labeling

CLEAR VISION EYE DROPS				
polyvinyl alcohol, povidone solution/ drops				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67751-212	
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	5 mg in 1 mL	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE, UNSPECIFIED - UNII:FZ989GH94E)		POVIDONE, UNSPECIFIED	6 mg in 1 mL	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0K00R)				
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
DEXTROSE (UNII: IY9XDZ35W2)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67751-212-01	1 in 1 CARTON	07/14/2021	
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	07/14/2021	

Labeler - Navajo Manufacturing Company Inc. (091917799)

Establishment			
Name	Address	ID/FEI	Business Operations
Navajo Manufacturing Company Inc.		136941411	relabel(67751-212) , repack(67751-212)
Establishment			

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(67751-212) , label(67751-212)

Revised: 7/2021

Navajo Manufacturing Company Inc.