POLYVINYL ALCOHOL- polyvinyl alcohol solution/ drops
AvPAK

Polyvinyl Alcohol Ophthalmic Solution

Active ingredient

Polyvinyl Alcohol 1.4%

Purpose

Lubricant

Uses

- for the temporary relief of burning, irritation and discomfort due to dryness of the eye or exposure to wind or sun
- may be used as a protectant against further irritation

Warnings

- **Do not use** if solution changes color or becomes cloudy

When using this product

- Avoid contamination, do not touch tip of container to any surface.
- Replace cap after use

Stop use and ask a doctor if

- you experience eye pain, changes in vision, continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222)

Directions

- Shake well before use
- instill 1 to 2 drops in the affected eye(s) as needed

Other information

- Store at room temperature 15-30C (59-86F)
- Do No Use if imprinted seal on cap is torn, broken or missing

- Discard 90 days after opening
- Retain outer carton for full product information

Inactive ingredients

dibasic sodium phosphate, edetate disodium, monobasic sodium phosphate,

purified water, sodium chloride. Phosphoric acid and/or sodium hydroxide may be added to adjust pH. PRESERVATIVE ADDED: benzalkonium chloride 0.01%

Questions ?

Call 1-855-361-3993

Distributed by:

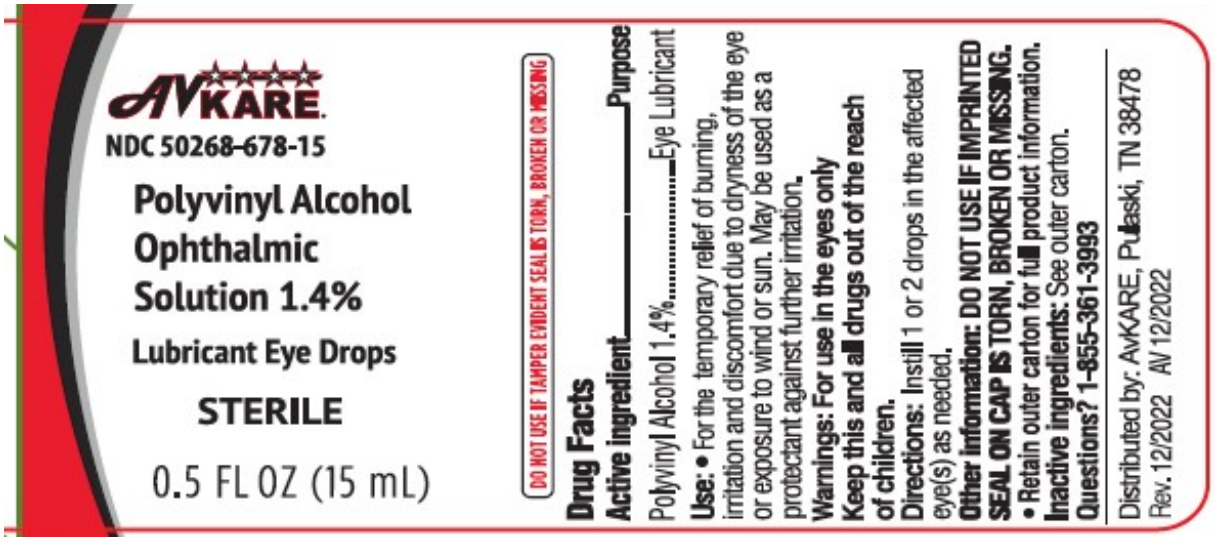
AvKARE

Pulaski, TN 38478

www.avkare.com

Rev. 12/2022 AV 12/2022

Package/Label Principal Display Panel



POLYVINYL ALCOHOL

polyvinyl alcohol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50268-678
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYVINYL ALCOHOL (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	POLYVINYL ALCOHOL	14 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50268-678-15	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/05/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	01/05/2023	

Labeler - AvPAK (832926666)

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