RUGBY LUBRICATING DROPS- polyvinyl alcohol solution/ drops
A-S Medication Solutions

Rugby Lubricating Eye Drops

Active ingredient

Polyvinyl Alcohol 1.4%

Purpose

Lubricant

Uses

- for temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun
- for use as a protectant against further irritation

Warnings

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

When using this product

- do not touch tip of container to any surface to avoid contamination
- remove contact lens before using.
- 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
- 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

Directions

- instill 1 to 2 drops in the affected eye(s) as needed

Other information

- store at room temperature
- keep tightly closed

- replace cap after use
- retain carton for full drug facts

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 1-800-645-2158

HOW SUPPLIED

Product: 50090-5996

NDC: 50090-5996-0 15 mL in a BOTTLE

polyvinyl alcohol solution/ drops

RUGBY LUBRICATING DROPS

polyvinyl alcohol solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-5996(NDC:0536-1325)
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
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-5996-0	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	04/19/2021	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-5996)

Revised: 11/2023 A-S Medication Solutions