RUGBY LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/drops

Rugby Laboratories

Rugby Lubricant Eye Drops 30ct and 50ct (PLD)

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Lubricant

Uses

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

Warnings

For external use only

Do not use this product if

solution changes color or becomes cloudy

When using this product

- do not reuse
- once opened, discard
- to avoid contamination, do not touch tip of container to any surface
- do not touch unit-dose tip to eye

Stop use and ask a doctor if

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

Keep out of reach of children.

If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

• to open, twist and pull tab to remove

- instill 1 or 2 drops in the affected eye(s) as needed and discard container
- if used for post-operative, (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions

Other information

- store at 15°-25°C (59°-77°F)
- use only if single-use container is intact
- use before expiration date marked on container
- RETAIN THIS CARTON FOR FUTURE REFERENCE

Inactive ingredients

calcium chloride, **hydrochloric acid, magnesium chloride, potassium chloride, purified water, sodium chloride, **sodium hydroxide, sodium lactate.

**May contain these ingredients to adjust pH.

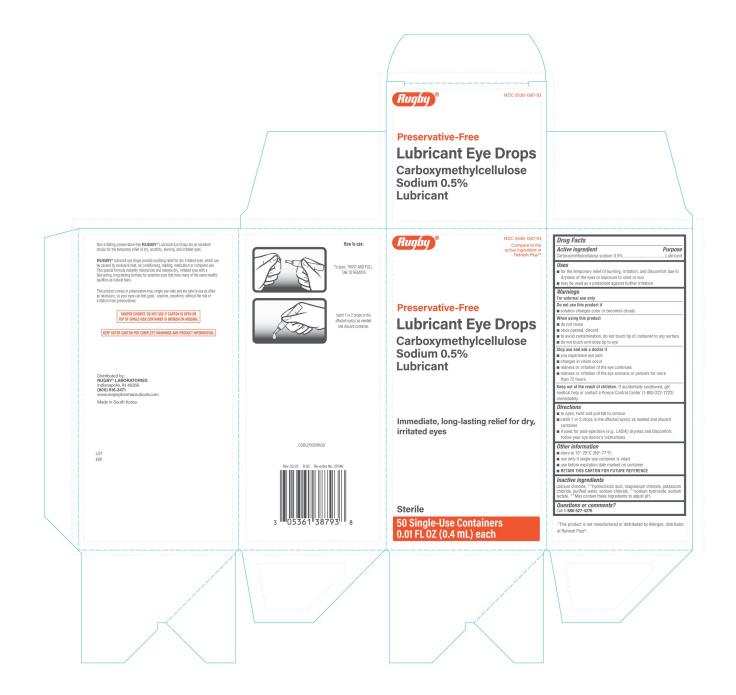
Questions or comments?

Call 1-888-527-4276

Rugby Lubricant Eye Drops 30ct



Rugby Lubricant Eye Drops 50ct



RUGBY LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0536-1387
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)				
POTASSIUM CHLORIDE (UNII: 660YQ98I10)				
HYDROCHLORIC ACID (UNII: QTT17582CB)				
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)				
WATER (UNII: 059QF0KO0R)				
SODIUM LACTATE (UNII: TU7HW0W0QT)				

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0536- 1387-92	30 in 1 BOX	03/23/2023	
1		0.4 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
2	NDC:0536- 1387-93	50 in 1 BOX	03/23/2023	
2		0.4 mL in 1 VIAL, SINGLE-USE; 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	03/23/2023	

Labeler - Rugby Laboratories (079246066)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(0536-1387) , label(0536-1387)

Establishment			
Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	manufacture(0536-1387), pack(0536-1387), label(0536-1387)

Revised: 12/2023 Rugby Laboratories