RUGBY CARBOXYMETHYLCELLULOSE SODIUM 0.5% EYE DROPS-
carboxymethylcellulose sodium solution/ drops
Rugby Laboratories

Rugby Carboxymethylcellulose Sodium 0.5% Eye Drops (PLD)

Active ingredient

Carboxymethylcellulose sodium 0.5%

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

For external use only

Do not use

- if this product changes color or becomes cloudy
- if you are sensitive to any ingredient in this product

When using this product

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

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 reach of children.

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

Directions

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

Other information

store at 15°-25°C (59°-77°F)

Inactive ingredients

benzalkonium chloride, boric acid, calcium chloride hydrate (dihydrate), hydrochloric acid**, magnesium chloride, potassium chloride, sodium borate, sodium chloride, sodium hydroxide**, water for injection

**May contain these ingredients to adjust pH.

Questions or comments?

Call 1-888-527-4276

Rugby Carboxymethylcellulose Sodium 0.5% Eye Drops 15mL

Rugby Carboxymethylcellulose Sodium 0.5% Eye Drops 2-15mL

RUGBY CARBOXYMETHYLCELLULOSE SODIUM 0.5% EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0536-1386
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0536-1386-94	1 in 1 BOX	05/11/2023	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:0536-1386-35	2 in 1 BOX	05/11/2023	
2		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	05/11/2023	

Labeler - Rugby Laboratories (079246066)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Revised: 12/2023 Rugby Laboratories