

PROEYE LUBE EYE DROPS- carboxymethylcellulose sodium 0.5 % solution/ drops

Renova Lifesciences Private Limited

PROEYE LUBE EYE DROPS

Active ingredients	Purpose
Carboxymethylcellulose Sodium 0.5 %	Lubricant

Uses

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

WARNINGS

- For external use only.
- To avoid contamination, do not touch tip of container to any surface. Replace cap after using.
- If solution changes color or becomes cloudy, do not use.

STOP USE

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

- Instill 1 or 2 drops in the affected eye(s) as needed.
- Use the solution within 1 month of opening the vial.
- Use before expiration date marked on container.
- **RETAIN THIS CARTON FOR FUTURE REFERENCE.**

STORAGE

Store at room temperature

INACTIVE INGREDIENTS:


Oxychloro Complex (As preservative), Glycerin, Calcium Chloride Dihydrate, Magnesium Chloride Hexahydrate, Boric Acid, Sodium Chloride, Potassium Chloride, Sodium hydroxide (to adjust pH), Purified Water.

QUESTIONS?

In the U.S. call toll-free 1-888-589-6879 (Mon-Fri 9AM-5PM MST)
info@proeye.in

www.proeye.in

- Instill 1 or 2 drops in the affected eye(s) as needed.
- For the temporary relief of burning, irritation and discomfort due to dryness of the eye or due to exposure to wind or sun.
- May be used as a protectant against further irritation.

Active Ingredients Carboxymethylcellulose Sodium 0.5%	Purpose Eye Lubricant	 NDC 83673-004-04	Warning : For external use only Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	Mfg. Lit. No. : G/28/1687
Inactive Ingredients Oxychloro Complex (As Preservative), Glycerin, Calcium Chloride Dihydrate, Magnesium Chloride Hexahydrate, Boric Acid, Sodium Chloride, Potassium Chloride, Sodium Hydroxide (to adjust pH), Purified Water.				Batch No. : Mfg. Date : Exp. Date :
Manufactured by : RENOVA LIFESCIENCES PRIVATE LIMITED Plot No : 35 / 36 / 37, R. K. Industrial Park, Phase - 1, Opp. : Bharat Benz Showroom, Off. Rajkot-Ahmedabad N.H. 8-B, Rampara - 360023, Dist - Rajkot, Gujarat - INDIA. TM - Trade Mark Applied for Proeye Lube		PROEYE LUBE lubricant eye drops <i>Immediate, soothing relief for dry eyes</i>		Directions : Instill 1 or 2 drops in the affected eye(s) as needed. Other information : Store at room temperature. Use the solution within 1 month of opening the vial. For directions and warnings kindly refer the carton. Questions? : In the U.S. call toll-free (888) 589-6879 (Mon-Fri 9AM-5PM MST) info@proeye.in www.proeye.in
		0.34 FL OZ (10 ml) Sterile		



PROEYE LUBE Carton

PROEYE LUBE Product Label

LUBRICANT EYE DROPS

Immediate, soothing relief for dry eyes

Lubricant Eye Drops instantly moisturizes and relieves dry, irritated eyes with a fast-acting, long-lasting formula that has many of the same healthy qualities as your own natural tears.

STERILE

One 10 mL Bottle (0.34 FL OZ)

RENOVA LIFESCIENCES PRIVATE LIMITED

Plot No: 35/36/37, R. K. Industrial Park, Phase - 1, Opp. Bharat Benz Showroom

Off. Rajkot-Ahmedabad N.H. 8-B, Rampara- 360023

Dist - Rajkot, Gujarat - INDIA

TM - Trade Mark Applied for Proeye Lube

Questions?:In the U.S. call toll-free (888) 589-6879

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FOR EXTERNAL USE ONLY

Uses:See carton for USES

WARNINGS:See carton for WARNINGS

PROEYE LUBE

PROEYE LUBE EYE DROPS

carboxymethylcellulose sodium 0.5 % solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83673-004
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORATE (UNII: T95DR77GMR)	

SODIUM CHLORITE (UNII: G538EBV4VF)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
BORIC ACID (UNII: R57ZHV85D4)	
GLYCERIN (UNII: PDC6A3C0OX)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CHLORINE DIOXIDE (UNII: 8061YMS4RM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83673-004-04	1 in 1 CARTON	08/01/2024	
1		10 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	08/01/2024	

Labeler - Renova Lifesciences Private Limited (873681128)

Establishment

Name	Address	ID/FEI	Business Operations
Renova Lifesciences Private Limited		956992404	manufacture(83673-004)

Revised: 8/2024

Renova Lifesciences Private Limited