

EYES ALIVE LUBRICATING- carboxymethylcellulose sodium, unspecified liquid
RIVIERA PHARMA INC

Eyes Alive™
Lubricating

Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Purpose

Eye 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 of the eye.

Warnings

For external use only

Do not use if solution changes color or becomes cloudy.

When using this product

- to avoid contamination, do not touch tip of container to any surface. Do not reuse. Once open, discard.
- do not touch unit dose tip to eye

Stop use and ask a doctor before use 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 right away.

Directions

- **TWIST AND REMOVE TAB**
- Place 1 or 2 drops in the affected eye(s) as needed for relief and discard container.
- If used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctors instructions.

Other information

- **Use only if single use container is intact.**
- **Use before expiration date marked on container.**
- **Store at 59°-86°F (15°-30°C)**
- **RETAIN THIS CARTON FOR FUTURE REFERENCE.**

Inactive ingredients

Calcium chloride, hydrochloric acid¹, magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide¹, and sodium lactate.

1 May or may not contain this ingredient to adjust pH.

Questions or comments?

800.477.2884, M-F 8AM-5PM Eastern Time

You can always report serious effects to this number.

PRINCIPAL DISPLAY PANEL - 0.6 mL Container Carton

NDC 83291-200-32

Eyes
Alive™

Lubricating Eye Drops
Carboxymethylcellulose sodium 0.5%

Preservative Free

IMMEDIATE RELIEF FOR
DRY IRRITATED EYES

32 Sterile Single-Use Containers
0.02 fl oz (0.6 mL) each

MADE IN THE USA

Preservative Free



Eyes Alive™ Lubricating Eye Drops

NDC 83291-200-32



Eyes Alive™ Lubricating Eye Drops Carboxymethylcellulose sodium 0.5% Preservative Free



IMMEDIATE RELIEF FOR
DRY IRRITATED EYES

32 Sterile Single-Use Containers
0.02 fl oz (0.6 mL) each
MADE IN THE USA

Eyes Alive™ Lubricating Eye Drops

Make your eyes feel **Alive** again with lubricating eye drops.

EyesAlive™ lubricating eye drops provide moisturizing relief and protection for dry, irritated, sensitive eyes.

EyesAlive™ drops come in preservative free, sterile, single use containers



MADE IN THE USA

Eyes Alive™ Lubricating Eye Drops

How to use:
TWIST AND REMOVE TAB to open. Place 1 or 2 drops in the affected eye(s) as needed for relief and discard container.



See Drug Facts panel for directions.



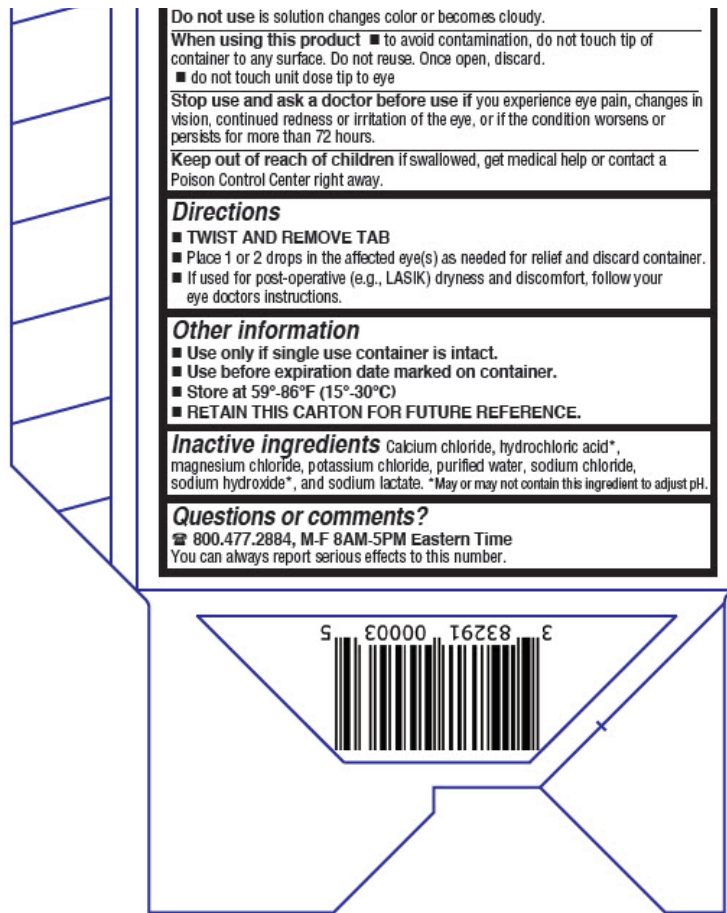
16100 SW Market Street
Indiantown, FL 34956

Drug Facts

Active Ingredients	Purpose
Carboxymethylcellulose sodium 0.5%	Eye 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 of the eye.

Warnings
For external use only



EYES ALIVE LUBRICATING

carboxymethylcellulose sodium, unspecified liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83291-200
Route of Administration	INTRAOCULAR		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE (UNII: M410D6VV5M)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

SODIUM LACTATE (UNII: TU7HW0W0QT)**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83291-200-02	2 in 1 POUCH	06/01/2023	
1		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
2	NDC:83291-200-04	4 in 1 POUCH	06/01/2023	
2		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
3	NDC:83291-200-32	32 in 1 CARTON	06/01/2023	
3		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
4	NDC:83291-200-52	52 in 1 CARTON	06/01/2023	
4		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
5	NDC:83291-200-72	72 in 1 CARTON	06/01/2023	
5		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
6	NDC:83291-200-00	100 in 1 CARTON	06/01/2023	
6		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
7	NDC:83291-200-08	8 in 1 CARTON	06/01/2023	
7		0.6 mL in 1 AMPULE; 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	06/01/2023	

Labeler - RIVIERA PHARMA INC (118748772)

Revised: 5/2024

RIVIERA PHARMA INC