EYES ALIVE LUBRICATING- carboxymethylcellulose sodium liquid
DIVISION 5 LABS, INC.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Eyes Alive
Lubricating

Drug Facts

Active ingredient (in each unit dose)

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 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 doctor's instructions.

Other Information

- **Use only if single-use container in intact.**
- **Use before expiration date marked on container.**
- **RETAIN THIS CARTON FOR FUTURE REFERENCE.**

Inactive ingredients

Calcium chloride, hydrochloric acid[1], magnesium chloride, potassium chloride, purified water, sodium chloride, sodium hydroxide[1], and sodium lactate.

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

Questions or comments?

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

You can also report serious side effects to this number.

PRINCIPAL DISPLAY PANEL - 32 Ampule Carton

NDC 69183-200-32

Eyes
***Alive*™**

Lubricating Eye Drops
Lubricant/carboxymethylcellulose sodium

Long lasting moisturizing
drops to keep your
Eyes Feeling Alive!

IMMEDIATE RELIEF FOR
DRY IRRITATED EYES

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

MADE IN THE USA

NDC 69183-200-32

Eyes Alive™

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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.

Division5 LABS. INC.™

Pharmaceutical Division

7555 West 2nd Court

Miami, FL 33014

DIVISION5LABS.COM

MADE IN THE USA

Lubricating Eye Drops

Eyes Alive™

Drug Facts

Active ingredient *(in each unit dose)*	***Purpose***
Carboxymethylcellulose sodium 0.5%	Eye lubricant

Uses

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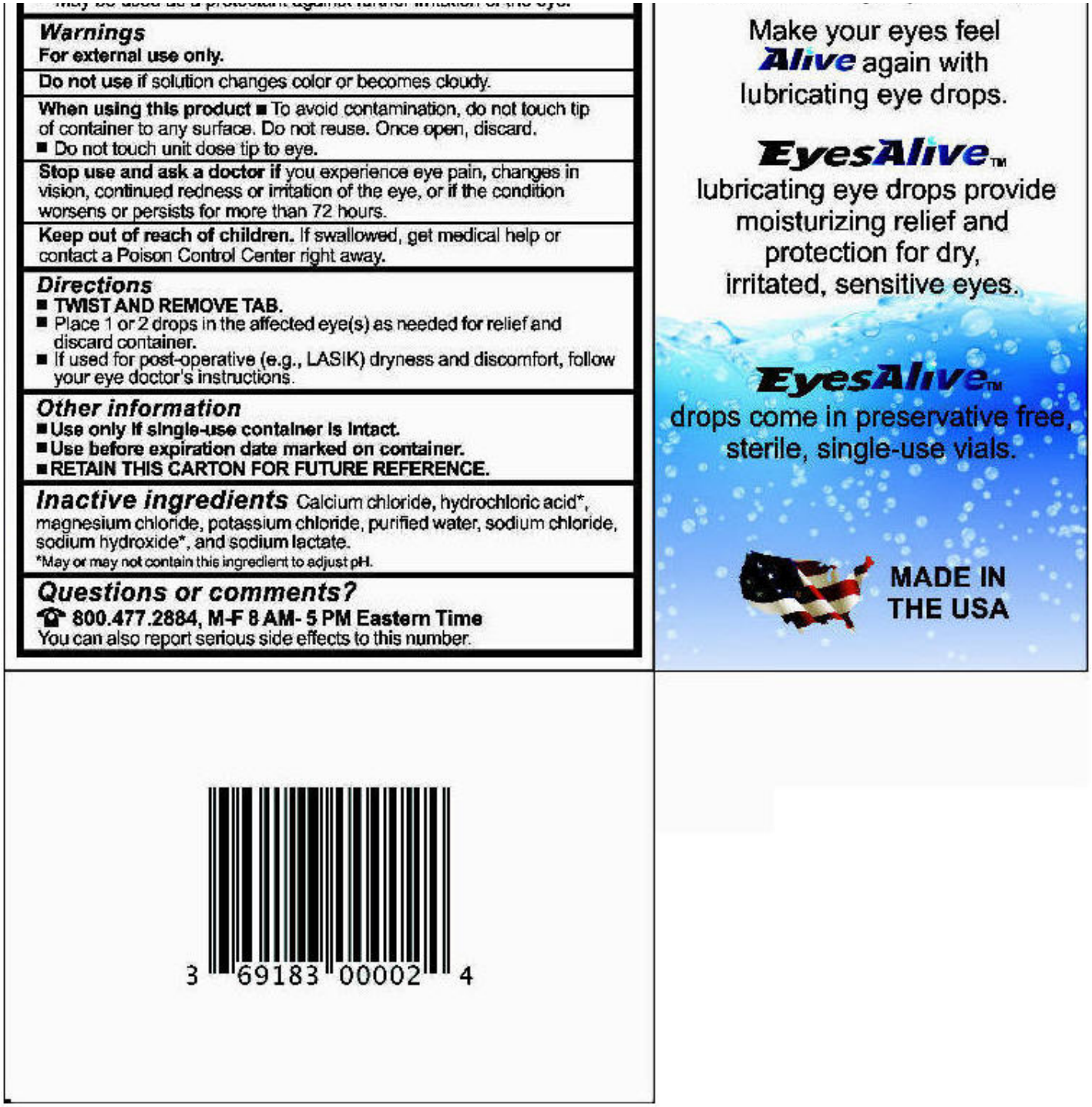
- May be used as a protectant against further irritation of the eye.

Eyes Alive™

Lubricating Eye Drops



EYES ALIVE LUBRICATING

carboxymethylcellulose sodium liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69183-200
Route of Administration	INTRAOCULAR		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
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:69183-200-02	2 in 1 POUCH	02/01/2015	
1		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
2	NDC:69183-200-04	4 in 1 POUCH	02/01/2015	
2		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
3	NDC:69183-200-32	32 in 1 CARTON	02/01/2015	
3		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
4	NDC:69183-200-52	52 in 1 CARTON	02/01/2015	
4		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
5	NDC:69183-200-72	72 in 1 CARTON	02/01/2015	
5		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
6	NDC:69183-200-00	100 in 1 CARTON	02/01/2015	
6		0.6 mL in 1 AMPULE; Type 0: Not a Combination Product		
7	NDC:69183-200-08	8 in 1 CARTON	02/01/2015	
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 FINAL	part349	02/01/2015	

Labeler - DIVISION 5 LABS, INC. (968198288)

Revised: 12/2019 DIVISION 5 LABS, INC.