EYES ALIVE EYECON LUBRICATING- carboxymethylcellulose sodium, unspecified form 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™ EYECON™
Lubricating

Drug Facts

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Eye Lubricant

Uses

- for the temporary relief of burning, irritation and discomfort due to dryness of the eye.
- 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 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 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

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

Other Information

- use before expiration date marked on container
- store at room temperature
- remove contact lenses before using
- 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?

1-800 477-2884 M-F 8 AM 5 PM Eastern Time

You can also report serious side effects to this number.

PRINCIPAL DISPLAY PANEL - 10 mL Bottle Carton

NDC 69183 205 01

Eyes
Alive™

EYECON™
LUBRICATING EYE DROPS
MULTI DOSE

FOR
DRY EYE
RELIEF

MAKE YOUR
EYES FEEL
Alive
AGAIN

USE AS
OFTEN AS
NEEDED

PRESERVATIVE FREE

STERILE
0.3 FL OZ (10 mL)

Over
240
Drops

MADE IN USA

EYES ALIVE EYECON LUBRICATING

carboxymethylcellulose sodium, unspecified form liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69183-205
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:69183-205-01	1 in 1 CARTON	01/07/2019	
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:69183-205-02	2 in 1 CARTON	01/07/2019	
2		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 final	part349	01/07/2019	

Labeler - DIVISION 5 LABS, INC. (968198288)

Revised: 11/2019 DIVISION 5 LABS, INC.