REFRESH OPTIVE MEGA-3 PF- carboxymethylcellulose sodium, glycerin, polysorbate 80 solution/ drops Allergan, Inc.

REFRESH OPTIVE MEGA-3 PF® Drug Facts

Active ingredients

Carboxymethylcellulose sodium 0.5%

Glycerin 1%

Polysorbate 80 0.5%

Purpose

Eye lubricant

Eye lubricant

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.

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, do not use.

Stop use and ask 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

- Prior to first use, please read the "Instructions for Use" inside this carton.
- Instill 1 or 2 drops in the affected eye(s) as needed.

Other information

- Use only if tape seals on top and bottom flaps are intact.
- Use before expiration date marked on container.
- Discard 90 days after opening.
- Store at 59°-77°F (15°-25°C).
- RETAIN THIS CARTON FOR FUTURE REFERENCE

Inactive ingredients

Boric acid; butylated hydroxyl toluene; carbomer copolymer type A; castor oil; erythritol; flaxseed oil; levocarnitine; polyoxyl 40 stearate; purified water; and trehalose. May contain hydrochloric acid and/or sodium hydroxide (to adjust pH).

Questions or comments?

1.800.678.1605

refresheyedrops.com

PRESERVATIVE-FREE MULTIDOSE BOTTLE

BUILT-IN DOUBLE LOCKOUT SYSTEM KEEPS DROPS STERILE.

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Product of US

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PRINCIPAL DISPLAY PANEL

NDC 0023-3988-10

NEW PRESERVATIVE-FREE

Refresh
OPTIVE
MEGA-3® PF
Lubricant Eye Drops

with Hydrocell®

Fast-Acting Dry Eye Relief

Enhanced With Flaxseed Oil*

0.33 fl oz (10 mL) Sterile



REFRESH OPTIVE MEGA-3 PF

carboxymethylcellulose sodium, glycerin, polysorbate 80 solution/ drops

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0023-3988

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety Ingredient Name Basis of Strength Strength CARBOXYMETHYLCELLULOSE SODIUM (UNII: K6790BS311) CARBOXYMETHYLCELLULOSE 5 mg (CARBOXYMETHYLCELLULOSE - UNII:05JZ17B19X) SODIUM in 1 mL 10 ma GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX) **GLYCERIN** in 1 mL POLYSORBATE 80 (UNII: 60ZP39ZG8H) (POLYSORBATE 80 -5 mg **POLYSORBATE 80** UNII:60ZP39ZG8H) in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
BORIC ACID (UNII: R57ZHV85D4)			
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)			
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)			
CASTOR OIL (UNII: D5340Y2I9G)			

ERYTHRITOL (UNII: RA96B954X6)	
LINSEED OIL (UNII: 84XB4DV00W)	
LEVOCARNITINE (UNII: 0G389FZZ9M)	
POLYOXYL STEARATE 40 (UNII: 13A4J4NH9I)	
WATER (UNII: 059QF0KO0R)	
TREHALOSE (UNII: B8WCK70T7I)	

Other Ingredients				
Ingredient Kind	Ingredient Name	Quantity		
May contain	HYDROCHLORIC ACID (UNII: QTT17582CB)			
May contain	SODIUM HYDROXIDE (UNII: 55X04QC32I)			

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:0023-3988- 10	1 in 1 CARTON	02/15/2025			
1		10 mL in 1 VIAL; 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	02/15/2025		

Labeler - Allergan, Inc. (144796497)

Revised: 6/2024 Allergan, Inc.