REFRESH TEARS- carboxymethylcellulose sodium solution/ drops
REFRESH TEARS- carboxymethylcellulose sodium
Allergan, Inc.

REFRESH TEARS®
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 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 or becomes cloudy, do not 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.

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.

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; calcium chloride dihydrate; magnesium chloride hexahydrate; potassium chloride; purified water; PURITE® (stabilized oxychloro complex); sodium borate decahydrate; and sodium chloride. May contain hydrochloric acid and/or sodium hydroxide (to adjust pH).

Questions or comments?

1.800.678.1605

refresheyedrops.com

v2.0DFL0798

PRINCIPAL DISPLAY PANEL

NDC 0023-0798-15
Refresh
TEARS®
Lubricant Eye Drops
Original Strength
Dry Eye Relief

0.5 fl oz (15 mL) Sterile



PRINCIPAL DISPLAY PANEL

NDC 0023-0795-65
5 BOTTLE MULTI-PACK

Refresh
TEARS®
Lubricant Eye Drops
Original Strength
Lubricates & Moisturizes
4 bottles 0.5 fl oz (15 mL) each
Plus 1 bottle 0.17 fl oz (5 mL) Sterile



REFRESH TEARS
carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0798
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	

SODIUM CHLORITE (UNII: G538EBV4VF)	
SODIUM CHLORATE (UNII: T95DR77GMR)	
CHLORINE DIOXIDE (UNII: 8061YMS4RM)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0798-04	1 in 1 CARTON	05/16/1997	
1		3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
2	NDC:0023-0798-03	1 in 1 CARTON	05/16/1997	
2		3 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
3	NDC:0023-0798-15	1 in 1 CARTON	05/16/1997	
3		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
4	NDC:0023-0798-01	2 in 1 CARTON	05/16/1997	
4		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		
5	NDC:0023-0798-02	4 in 1 CARTON	05/16/1997	01/27/2018
5		15 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	05/16/1997	

REFRESH TEARS

carboxymethylcellulose sodium kit

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0023-0795

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0795-	1 in 1 CARTON	01/01/2000	

1	65	1 in 1 CARTON	01/01/2000	
1		1 in 1 TRAY; Type 1: Convenience Kit of Co-Package		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	4 BOTTLE, DROPPER	60 mL in 4
Part 2	1 BOTTLE, DROPPER	5 mL

Part 1 of 2

REFRESH TEARS

carboxymethylcellulose sodium solution/ drops

Product Information	
Item Code (Source)	NDC:0023-0796
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORITE (UNII: G538EBV4VF)	
SODIUM CHLORATE (UNII: T95DR77GMR)	
CHLORINE DIOXIDE (UNII: 8061YMS4RM)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0023-	15 mL in 1 BOTTLE, DROPPER; Type 0: Not a		

1	0796-15	Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	01/01/2000	

Part 2 of 2

REFRESH TEARS

carboxymethylcellulose sodium solution/ drops

Product Information

Item Code (Source)	NDC:0023-0797
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORITE (UNII: G538EBV4VF)	
SODIUM CHLORATE (UNII: T95DR77GMR)	
CHLORINE DIOXIDE (UNII: 8061YMS4RM)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0023-0797-05	5 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	01/01/2000	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	01/01/2000	

Labeler - Allergan, Inc. (144796497)

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