

**SIGNATURE CARE LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/
drops**

Better Living Brands LLC

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.

Active ingredient

Carboxymethylcellulose sodium 0.5%

Purpose

Carboxymethylcellulose sodium.....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.

⚠ Do not use this product if

- solution changes color or becomes cloudy

When using the product

- do not reuse
- once opened, discard
- to avoid contamination, do not touch tip of container to any surface
- do not touch unit-dose tip to eye

Stop use and ask a doctor if

- you experience eye pain
- changes in vision occur
- redness or irritation of the eye continues
- redness or irritation of the eye worsens or persists for more than 72 hours

Keep out of reach of children. If accidentally swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

Directions

- to open, twist and pull tab to remove.
- instill 1 or 2 drops in the affected eye(s) as needed and discard container
- if used for post-operative (e.g., LASIK) dryness and discomfort, follow your eye doctor's instructions.

Other information

- store at 15°-25°C (59°-77°F).
- use only if single-use container is intact
- use before expiration date marked on container.
- **RETAIN THIS CARTON FOR FUTURE REFERENCE**

Inactive ingredients

calcium chloride, magnesium chloride, potassium chloride, purified water, sodium

chloride, and sodium lactate. May contain sodium hydroxide and/or hydrochloric acid to adjust pH.

Drug Facts

Active Ingredient Carboxymethylcellulose sodium 0.5% **Purpose**
Lubricant

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How to use:



To open, TWIST AND PULL TAB TO REMOVE



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*This product is not manufactured or distributed by Allergan, distributor of Refresh Plus®. Refresh Plus® is a registered trademark of Allergan.



Single Use | Preservative Free
Lubricant
Eye Drops

Signature Care
Quality Guaranteed

STERILE

Single Use | Preservative Free
Lubricant
Eye Drops

30 sterile dispensers Immediate, long-lasting relief for dry, irritated eyes

30 - 0.01 FL OZ (0.4 mL) DISPENSERS
0.01 FL OZ (0.4 mL) EA

Compare to Refresh Plus® active ingredient*

Signature Care
Quality Guaranteed

Non-irritating, preservative-free Signature Care™ Lubricant Eye Drops are an excellent choice for the temporary relief of dry, scratchy, burning, and irritated eyes.

Signature Care™ lubricant eye drops provide soothing relief for dry irritated eyes, which can be caused by excessive heat, air conditioning, reading, medication or computer use. This special formula instantly moisturizes and relieves dry, irritated eyes with a fast-acting, long-lasting formula for sensitive eyes that have many of the same healthy quality as natural tears.

This product comes in preservative-free, single use vials and are safe to use as often as necessary, so your eyes can feel good - anytime, anywhere, without the risk of irritation from preservatives.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPEN OR TOP OF SINGLE-USE CONTAINER IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY
BETTER LIVING BRANDS LLC
P.O. BOX 99
PLEASANTON, CA 94566-0099
1-888-723-3929
www.betterlivingbrandsLLC.com

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SIGNATURE CARE LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-705
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311) (CARBOXYMETHYLCELLULOSE - UNII:05JZI7B19X)	CARBOXYMETHYLCELLULOSE SODIUM	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	

POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-705-01	30 in 1 CARTON	04/11/2019	
1		0.4 mL in 1 VIAL, DISPENSING; 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	04/11/2019	

Labeler - Better Living Brands LLC (009137209)

Registrant - Unimed Pharmaceuticals, Inc. (689852052)

Establishment			
Name	Address	ID/FEI	Business Operations
Unimed Pharmaceuticals, Inc.		689852052	label(21130-705) , manufacture(21130-705) , pack(21130-705)

Revised: 4/2019

Better Living Brands LLC