UP AND UP DRY EYE RELIEF LUBRICANT EYE DROPS- carboxymethylcellulose sodium solution/ drops
Hanlim

Up & Up Dry Eye Relief Lubricant Eye Drops- Twin Pack (MFG Listing for PLD)

Carboxymethylcellulose Sodium 0.5%

Purpose.....Lubricant

If swallowed, get medical help or contact a Poison Control Center immediately (1-800-222-1222) right away.

Instill 1 or 2 drops in the affected eye(s) as needed

- For the temporary relief of burning, irritation, and discomfort due to dryness of the eye or from irritation from wind or sun.
- May be used to protect against further irritation.

WARNINGS
- Warnings - For external use only.

DO NOT USE
- Do not use - if solution changes color or becomes cloudy
- If you are sensitive or allergic to any ingredient in this product

WHEN USING
- To avoid contamination, do not touch tip of container to any surface.
- Replace cap after using.

STOP USE
Stop use and ask a doctor if
- you experience eye pain
- changes in vision occur
- redness or irritation of the eye(s) gets worse or lasts more than 72 hours

Inactive ingredients: benzalkonium chloride, boric acid, calcium chloride hydrate, hydrochloric acid, magnesium chloride, potassium chloride, sodium borate, sodium chloride, sodium hydroxide, water

UP AND UP DRY EYE RELIEF LUBRICANT EYE DROPS

carboxymethylcellulose sodium solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11716-0771
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
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11716-0771-1	2 in 1 BOX	10/31/2025	
1		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	10/31/2025	

Labeler - Hanlim (687986034)

Revised: 10/2025 Hanlim