DRIEYES LUBRICANT EYE DROPS- carboxymethylcellulose sodium 0.50% lubricant eye drops liquid
Jiangxi Hemei Pharmaceutical Co., Ltd

84010-095

Active Ingredient

Carboxymethylcellulose sodium 0.50%

Purpose

Lubricant

Use

·For use as a protectant against further irritation or to relieve dryness ofthe eye.

·For the temporary relief of burning & irritation due to the dryness of the eye.

·Relieves redness of the eye due to minor eye irritations.

Warnings

For external use only.

Do not use

if solution changes color or becomes cloudy.

When Using

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

Stop Use

·you experience eye pain
·you experience changes in vision
·you experience continued redness orirritation of the eye
·the condition worsens
·symptoms last for more than 72 hours

Ask Doctor

if you have narrow angle glaucoma

Keep Oot Of Reach Of Children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Instill 1 to 2 drops in the affected eye(s) up to four times daily.

Other information

· Store at room temperature.
· Remove contact lenses before using.

·Do not use if neckband on bottle is broken or missing

Inactive ingredients

Water□Sodium Hyaluronate□Boric acid□Borax□Borneol□Methylparaben□Ethanol

PRINCIPAL DISPLAY PANEL

DRIEYES LUBRICANT EYE DROPS

carboxymethylcellulose sodium 0.50% lubricant eye drops liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84010-095
Route of Administration	INTRAOCULAR		

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**
WATER (UNII: 059QF0KO0R)	
BORIC ACID (UNII: R57ZHV85D4)	
ALCOHOL (UNII: 3K9958V90M)	

METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM HYALURONATE (UNII: YSE9PPT4TH)	
BORNEOL (UNII: M89NIB437X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84010-095-01	15 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	03/28/2025	

Labeler - Jiangxi Hemei Pharmaceutical Co., Ltd (724892056)

Establishment

Name	Address	ID/FEI	Business Operations
Jiangxi Hemei Pharmaceutical Co., Ltd		724892056	manufacture(84010-095)

Revised: 3/2025 Jiangxi Hemei Pharmaceutical Co., Ltd