RETAINE HPMC- hypromellose 2910 (4000 mpa.s) solution/ drops
OCuSOFT, Inc.

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.

Drug Facts

Active Ingredient
Hypromellose 0.3%

Purpose
Eye lubricant

Uses
▪ For the temporary releif 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 ittitation.

Warnings
▪ For use in the eyes only.

Do not use if solution changes color or becomes cloudy.

When using this product
▪ do not touch tip of container to any surface to avoid contamination.
▪ replace cap after use.

Stop use and ask a doctor if
▪ you experience eye pain, changes in vision, continued redness or irritation of the eye.
▪ 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 immediately.

Directions

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

Other Information

▪ Use only it tamper seals on top ad bottom flaps are intact.
▪ Store between 15°-30°C (59°-86°F).
▪ Keep carton for complete product information.
▪ Discard after 12 weeks.

Inactive ingredients

Citric acid, sodium citrate, sorbitol, water for injection.

Questions?

(800)233-5469 M-F
8:30AM-5:00PM CST
www.retainebrand.com

Principal Display Panel

RETAINE HPMC

hypromellose 2910 (4000 mpa.s) solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54799-896
Route of Administration	OPHTHALMIC, TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Hypromellose 2910 (4000 Mpa.s) (UNII: RN3152OP35) (Hypromellose 2910 (4000 Mpa.s) - UNII:RN3152OP35)	Hypromellose 2910 (4000 Mpa.s)	3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Anhydrous Citric Acid (UNII: XF417D3PSL)	
Sodium Citrate (UNII: 1Q73Q2JULR)	
Sorbitol (UNII: 506T60A25R)	
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54799-896-10	10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	05/01/2013	

Labeler - OCuSOFT, Inc. (174939207)

Registrant - OCuSOFT, Inc. (174939207)

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