EYELLIC EYE DROPS (STERILE)- eye drops solution/ drops
Hi-Healthcare Inc

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Active Ingredients

Hypromelloses 0.2%
Polysorbate 80 0.5%
Naphazoline Hydrochloride 0.025%
Zinc Sulfate 0.25%

Purpose

Hypromelloses Lubricant
Polysorbate 80 Lubricant
Naphazoline Hydrochloride Redness Reliever
Zinc Sulfate Astringent

Uses

- For the temporary relief of discomfort and redness of the eye due to minor eye irritations.
- For the temporary relief of burning & irritation due to dryness of the eye.
- For protection against further irritation.

Directions

• Instill 1 or 2 drops in the affected eye(s) up to four times daily.
• Children under 6 years of age, ask a doctor.

Warnings

• For external use only.
• Do not use if solution changes color or becomes cloudy.
• Ask a doctor before use if you have narrow angle glaucoma.

When using this product

• Pupils may become enlarged temporarily.
• To avoid contamination, do not touch tip of container to any surface. Replace cap after using.
• Overuse may produce increased redness of the eye.
• Remove contact lens before using.

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.

If pregnant or breast-feeding, ask a health professional before use.

Other information

• Some users may experience initial stinging sensation.
• Store at 15° to 25°C (59° to 77°F).
• Keep tightly closed.
• Use before expiration date marked on the carton and bottle.

Questions or Comments?

Contact Hi-Healthcare, Inc.
1-877-578-6288
www.hi-healthcare.com

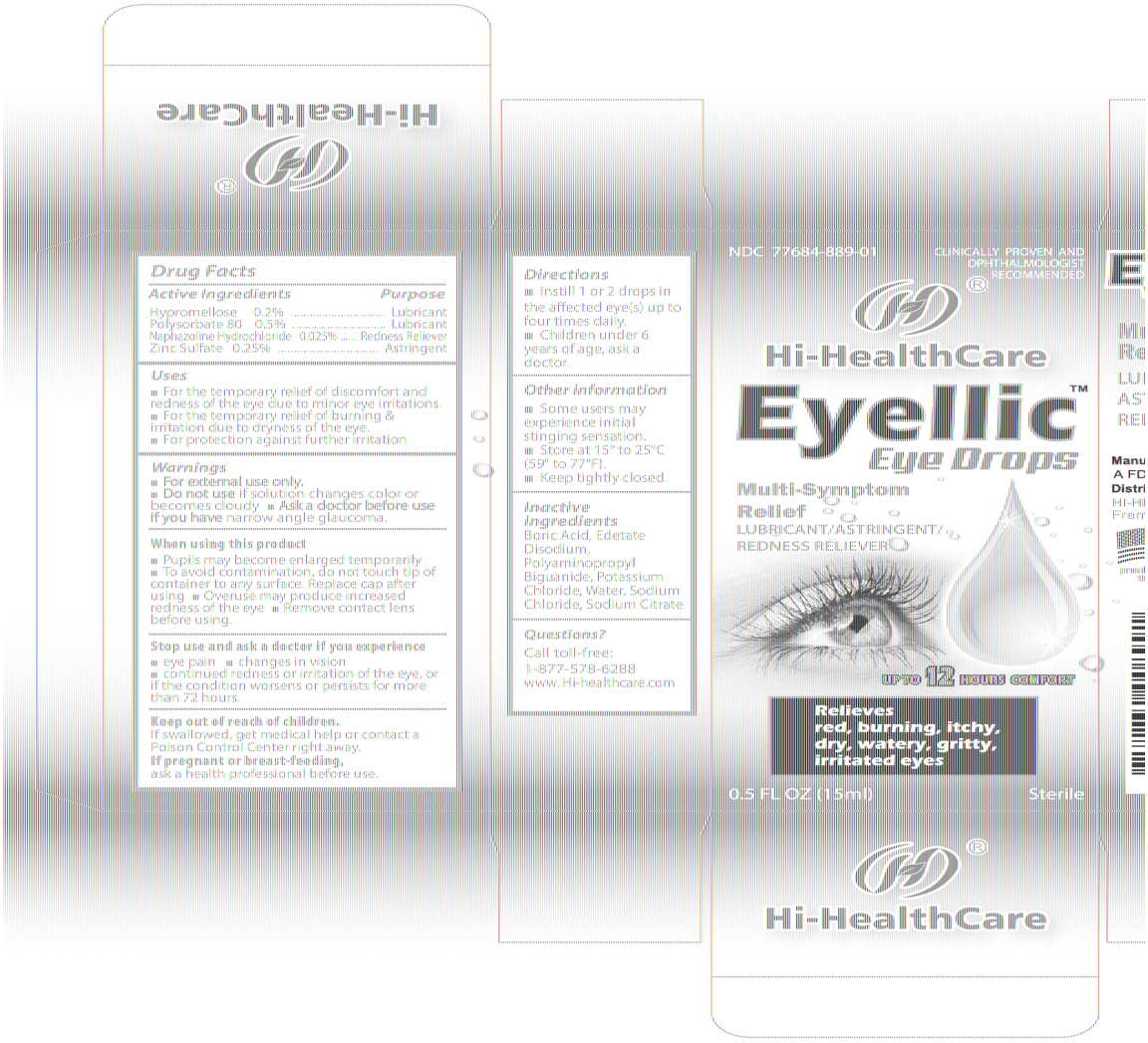
Distributed by Hi-Healthcare Inc.

Made in USA

Inactive ingredients

Boric Acid, Edetate Disodium, Polyaminopropyl Biguanide, Potassium Chloride, Water, Sodium Chloride, Sodium Citrate

Drug Facts Label



EYELLIC EYE DROPS (STERILE)

eye drops solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77684-899
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYPROMELLOSES (UNII: 3NXW29V3WO) (HYPROMELLOSES - UNII:3NXW29V3WO)	HYPROMELLOSES	0.2 g in 100 mL
POLYSORBATE 80 (UNII: 6OZP39ZG8H) (POLYSORBATE 80 - UNII:6OZP39ZG8H)	POLYSORBATE 80	0.5 g in 100 mL
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.025 g in 100 mL
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	0.25 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLYAMINOPROPYL BIGUANIDE HYDROCHLORIDE (UNII: 6P1MZW6VKY)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77684-899-01	15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/22/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
export only		05/22/2020	

Labeler - Hi-Healthcare Inc (087818745)

Revised: 5/2020 Hi-Healthcare Inc