# GOODSENSE IRRITATION RELIEF EYE- tetrahydrozoline hydrochloride, and zinc sulfate solution/ drops HANLIM PHARM. CO., LTD.

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.

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Active ingredients Purpose

Zinc Sulfate...... Astringent

#### Uses

- for the temporary relief of redness and irritation of the eye and for use as a protectant against further irritation.
- for the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun.

## Warnings

Ask a doctor before use if you have narrow angle glaucoma

When using this product

- pupils may become enlarged temporarily
- overuse may cause more eye redness
- remove contact lenses before using
- do not use if this solution change color or becomes cloudy
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye persists
- condition worsens or lasts more than 72 hours

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

Keep out of the reach of children.

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

#### Directions

- To open bottle, push cap down and twist counterclockwise. To close bottle, twist clockwise until it stops turning
- Put 1 to 2 drops in the affected eye(s) up to 4 times daily
- Children under 6 years of age: ask a doctor

### Other information

- some users may experience a brief tingling
- store at 15° to 25°C (59° to 77°F)

Inactive Ingredients: Benzalkonium Chloride, Boric Acid, Edetate Disodium, Purified Water, Sodium Chloride, Sodium Citrate

Distributed By:

Geiss, Destin & Dunn, Inc. Peachtree City, GA 30269 www.valuelabels.com 1-866-696-0957 Made in Korea



### GOODSENSE IRRITATION RELIEF EYE

tetrahydrozoline hydrochloride, and zinc sulfate solution/ drops

### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11716-0003
Route of Administration	ОРНТНАЬМІС		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
TETRAHYDRO ZOLINE HYDRO CHLO RIDE (UNII: 0 YZT43HS7D) (TETRAHYDRO ZOLINE - UNII:S9 U0 25 Y0 77)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL		
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	2.5 mg in 1 mL		

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
BORIC ACID (UNII: R57ZHV85D4)			
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)			
WATER (UNII: 059QF0KO0R)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:11716-0003-8	1 in 1 CARTON				
1		15 mL in 1 BOTTLE				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	08/02/2010		

# Labeler - HANLIM PHARM. CO., LTD. (687986034)

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