

**SHOPKO RELIEF - tetrahydrozoline hydrochloride and zinc sulfate solution**  
**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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**Drug Facts**

Active Ingredients	Purpose
Tetrahydrozoline HCL 0.05% .....	Redness Reliever
Zinc Sulfate 0.25% .....	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 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 changes color or become 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 lasts
- 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**

- Instill 1 or 2 drops in the affected eye(s) up to 4 times daily
- Store at 15° to 25°C (59° to 77°F)
- Children under 6 years of age: Ask a doctor

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

Distributed by

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703

www.ueccorp.com



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## SHOPKO RELIEF

tetrahydrozoline hydrochloride and zinc sulfate solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11716-0460
Route of Administration	OPHTHALMIC		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0 YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC - UNII:J41CSQ7QDS)	ZINC SULFATE	2.5 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11716-0460-1	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	10/08/2010	

**Labeler** - HANLIM PHARM. CO., LTD. (687986034)**Registrant** - UNITED EXCHANGE CORP. (840130579)**Establishment**

Name	Address	ID/FEI	Business Operations
HANLIM PHARM. CO., LTD.		687986034	manufacture

Revised: 10/2010

HANLIM PHARM. CO., LTD.