WALGREENS ORIGINAL EYE DROPS - tetrahydrozoline hydrochloride 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.

Drug Facts

Active ingredient

Purpose

Tetrahydrozoline HCL 0.05%......Redness reliever

Uses

• for the relief of redness of the eyes due to minor eye irritations

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 changes color or becomes cloudy
- to avoid contamination, do not touch tip
- 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 reach of children.

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

Directions

- 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 sensation
- store at 15° to 25°C (59° to 77°F)

Inactive ingredients: benzalkonium chloride, boric acid, edetate disodium, purified water, sodium chloride, and sodium borate



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WALGREENS ORIGINAL EYE DROPS

tetrahydrozoline hydrochloride solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11716-1168	
Route of Administration	OPHTHALMIC			

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	

Inactive Ingredients		
Ingredient Name	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
BORIC ACID (UNII: R57ZHV85D4)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
WATER (UNII: 059QF0KO0R)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SODIUM BORATE (UNII: 91MBZ8H3QO)		

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:11716-1168-6	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	05/26/2010		

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

Revised: 5/2010 HANLIM PHARM. CO., LTD.