GOODSENSE EYE DROPS ORIGINAL FORMULA- tetrahydrozoline hcl solution
Geiss, Destin & Dunn, Inc.

GoodSense Eye Drops Original (PLD)

Active ingredient

Tetrahydrozoline HCl 0.05%

Purpose

Tetrahydrozoline HCl ..... Redness reliever

Use

- relieves redness of the eye due to minor eye irritation

Warnings

For external use only

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.
- if solution changes color or becomes cloudy, do not use
- 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

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 [1-800-222-1222] right away.

Directions

Instill 1 to 2 drops in the affected eye(s) up to 4 times daily

Other information

store at 15º-30ºC (59º-86ºF)

Inactive ingredients

benzolkonium chloride, boric acid, edetate disodium, purified water, sodium borate, sodium chloride

GOODSENSE EYE DROPS ORIGINAL FORMULA

tetrahydrozoline hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-141
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-141-01	1 in 1 CARTON	01/03/2020	
1		15 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	01/03/2020	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

Registrant - K.C. Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(50804-141) , pack(50804-141) , label(50804-141)

Revised: 12/2023 Geiss, Destin & Dunn, Inc.