MEDICS CHOICE EYE DROPS ORIGINAL REDNESS RELIEVER- tetrahydrozoline hydrochloride solution/ drops
KC Pharmaceuticals, Inc.

Medic's Choice Eye Drops Original Redness Reliever

Active ingredient

Tetrahydrozoline HCl 0.05%

Purpose

Tetrahydrozoline HCl.....Redness reliever

Use

■ relieves redness of the eye due to minor eye irritations

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 15°-30°C (59°-86°F)

Inactive ingredients

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

MEDICS CHOICE EYE DROPS ORIGINAL REDNESS RELIEVER

tetrahydrozoline hydrochloride solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55651-005
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-005-01	1 in 1 CARTON	08/28/2003	
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	08/28/2003	

Labeler - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(55651-005) , pack(55651-005) , label(55651-005)

Revised: 1/2024 KC Pharmaceuticals, Inc.