

**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

Medic's Choice
EYE DROPS ORIGINAL

Drug Facts
Active ingredient Tetrahydrozoline HCl 0.05%
Purpose 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

Drug Facts (continued)
Other information
store at 15°-30°C (59°-86°F)

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

*This product is not manufactured or distributed by Johnson & Johnson, Healthcare Products, distributor of Visine® Original.

Medic's Choice
EYE DROPS ORIGINAL
Redness Reliever
Fast acting formula
Sterile
0.5 fl. oz. (15mL)

Compare to the active ingredient in Visine® Original*

NDC 55651-005-01

KC Pharmaceuticals, Inc.
Pomona, CA 91768
Questions? Call 1-888-527-4276
Made in USA

TAMPER EVIDENT: DO NOT USE IF IMPRINTED NECKBAND IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

LOT
EXP

0 95072 00551 0
CEDRIG0051MCT2

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.