QUALITY CHOICE EYE DROPS IRRITATION RELIEF- tetrahydrozoline hcl, zinc sulfate solution/ drops
Quality Choice

Quality Choice Eye Drops Irritation Relief (PLD)

Active ingredients

Tetrahydrozoline HCl 0.05%

Zinc sulfate 0.25%

Purposes

Redness reliever

Astringent

Use

- for temporary relief of discomfort and 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 lenses 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 or 2 drops in the affected eye(s) up to 4 times daily.

Other information

store at 15º-30ºC (59º-86ºF)

Inactive ingredients

benzalkonium chloride, boric acid, edetate disodium, purified water, sodium chloride, sodium citrate.

Quality Choice Eye Drops Irritation Relief 15mL

QUALITY CHOICE EYE DROPS IRRITATION RELIEF

tetrahydrozoline hcl, zinc sulfate solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-190
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
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	0.25 g

ZINC SULFATE (UNII: 89D30H96TB) (ZINC CATION - UNII:13S1S83F37)	ZINC CATION	in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-190-14	1 in 1 BOX	08/11/2024	
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	08/11/2024	

Labeler - Quality Choice (011920774)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(83324-190) , pack(83324-190) , label(83324-190)

Revised: 8/2024 Quality Choice