QUALITY CHOICE EYE DROPS IRRITATION RELIEF- tetrahydroxoline hcl, zinc sulfate solution
Quality Choice

Quality Choice Eye Drops Irritation Relief (PLD)

Active ingredients

Tetrahydrozoline HCl……0.05%

Zinc sulfate……0.25%

Purpose

Tetrahydrozoline HCl……Redness reliever

Zinc sulfate……Astringent

Uses

- for temporarry 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 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 right away.

Directions

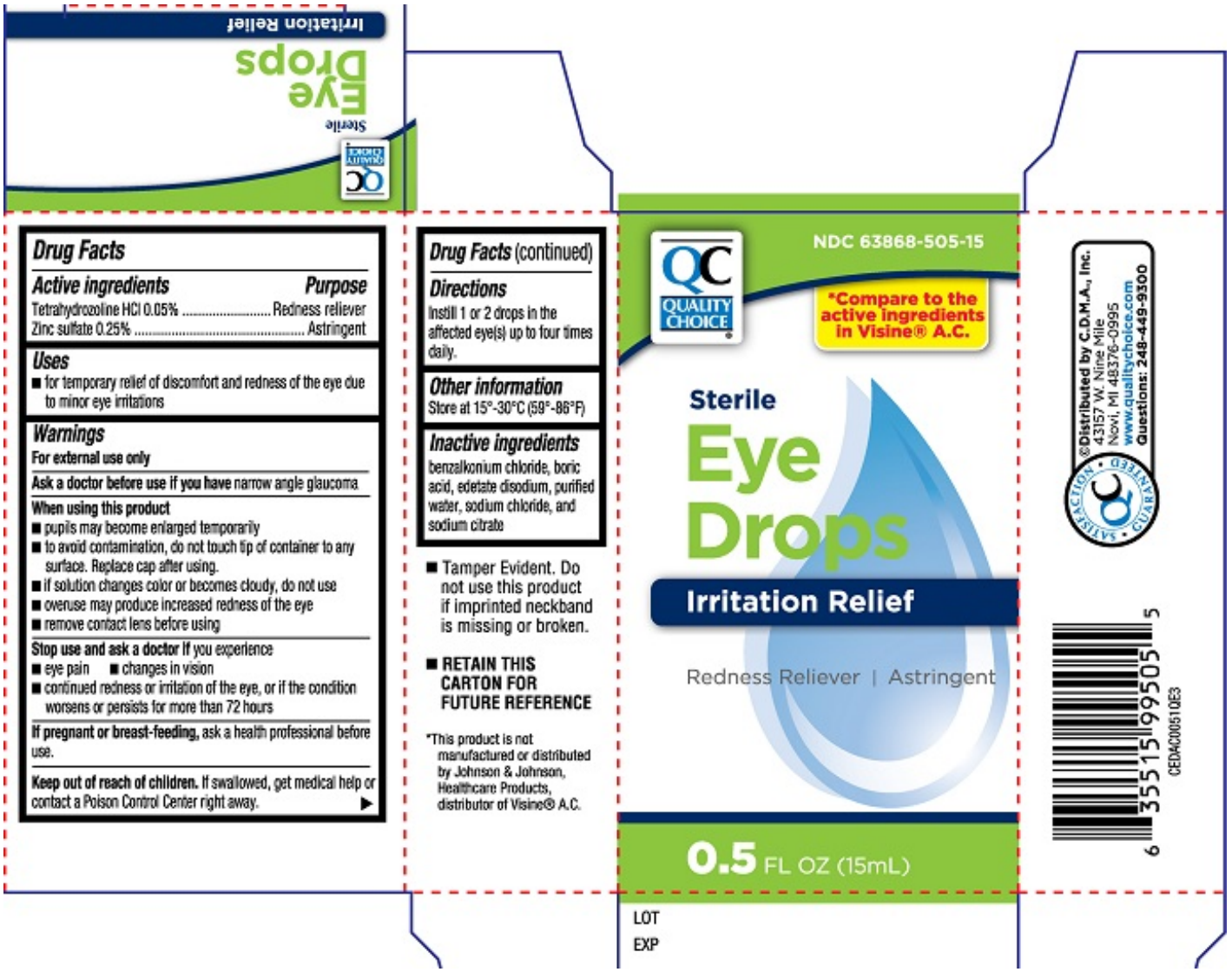
Instill 1 or 2 drops in the affected eye(s) up to four times daily.

Other information

store at 15º-30ºC (59º-86ºF)

Inactive ingredients

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



QUALITY CHOICE EYE DROPS IRRITATION RELIEF

tetrahydroxoline hcl, zinc sulfate solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-505
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 in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
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:63868-505-15	1 in 1 CARTON	01/05/2020	
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	01/05/2020	

Labeler - Quality Choice (011920774)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(63868-505) , pack(63868-505) , label(63868-505)

Revised: 12/2023 Quality Choice