QUALITY CHOICE EYE DROPS REDNESS RELIEF- tetrahydrozoline hcl solution/ drops
Chain Drug Marketing Assoc., Inc.

Quality Choice Eye Drops Redness Relief 15mL (PLD)

Active ingredient

Tetrahydrozoline HCL 0.05%

Purpose

Redness reliever

Uses

- 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 chlidren

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 four times daily

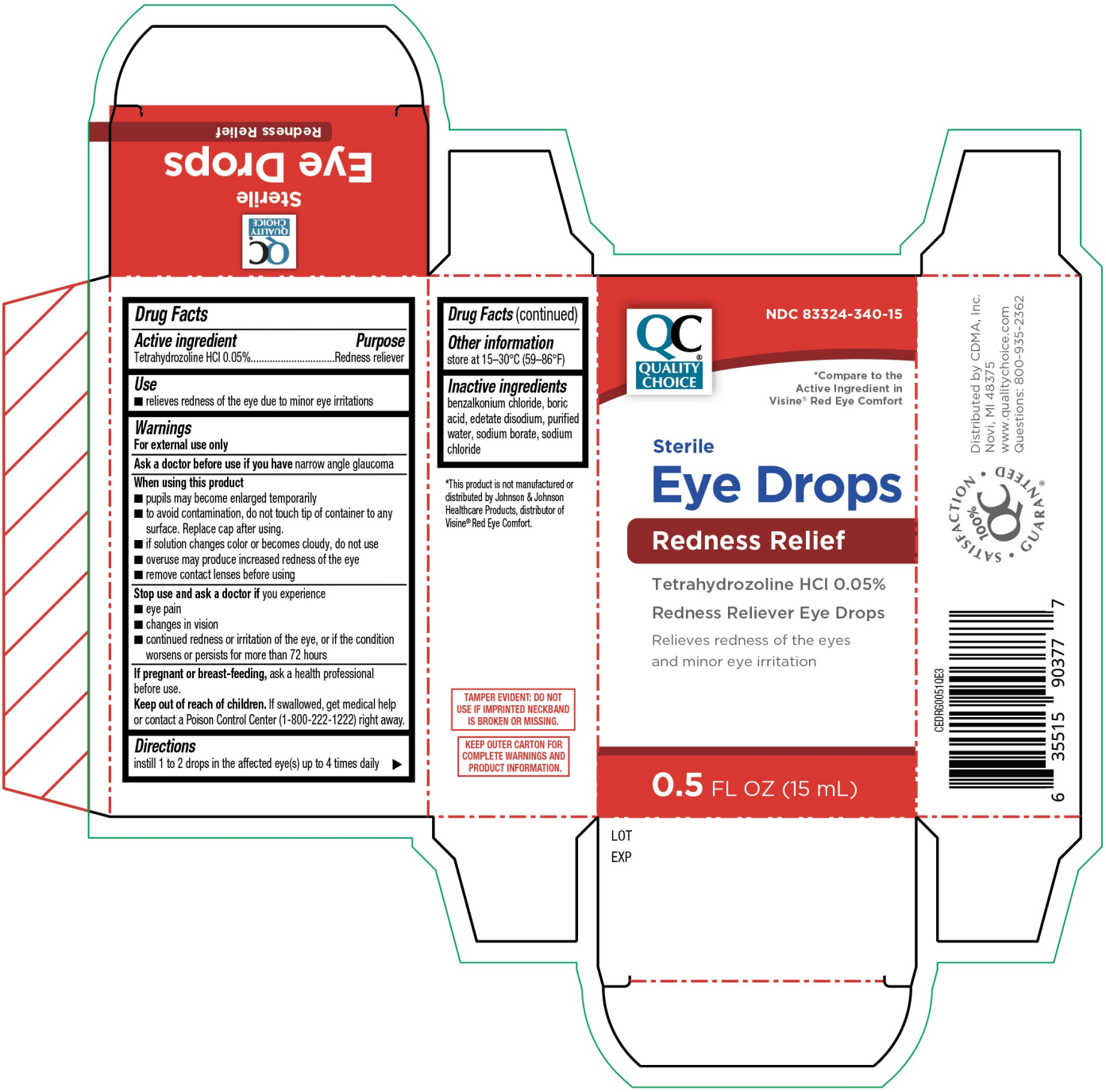
Other information

store at 15-30°C (59°-86°F)

Inactive ingredients

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

Quality Choice Eye Drops Redness Relief 15mL



QUALITY CHOICE EYE DROPS REDNESS RELIEF

tetrahydrozoline hcl solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-340
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
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-340-15	1 in 1 CARTON	06/26/2025	
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	06/26/2025	

Labeler - Chain Drug Marketing Assoc., Inc. (011920774)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(83324-340) , pack(83324-340) , label(83324-340)

Revised: 6/2025 Chain Drug Marketing Assoc., Inc.