EQUALINE EYE DROPS A.C.- tetrahydrozoline hcl, zinc sulfate solution
United Natural Foods, Inc.

Equaline Eye Drops A.C.15 mL (PLD)

Active ingredients

Tetrahydrozoline HCl....0.05%

Zinc sulfate.....0.25%

Purposes

Tetrahydrozoline HCl.....Redness reliever

Zinc sulfate.....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 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 (800-222-1222) right away.

Directions

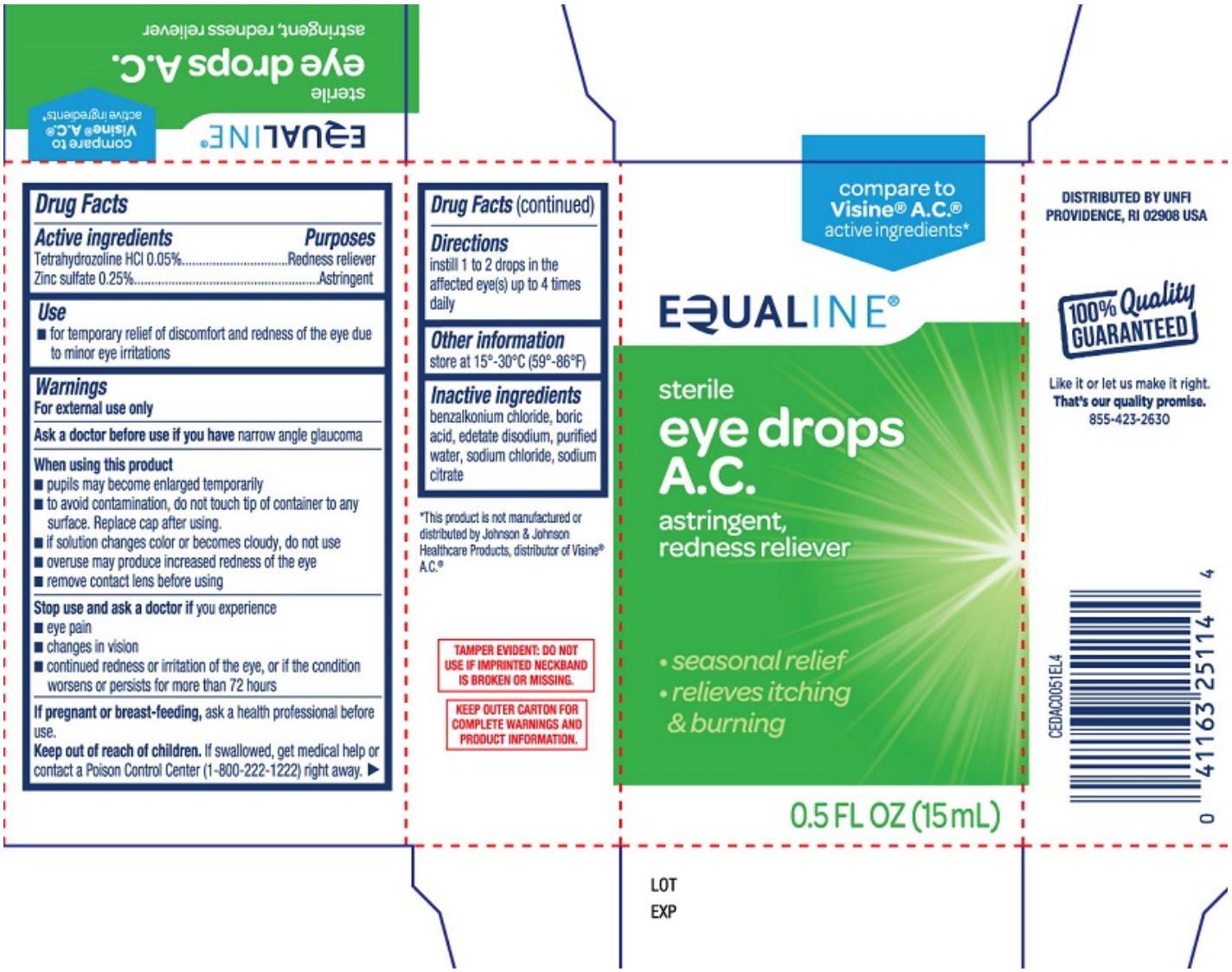
Instill 1 to 2 drops in the affected eye(s) up to 4 times daily.

Other information

store at 15º-30ºC (59º-86ºF)

Inactive ingredients

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



EQUALINE EYE DROPS A.C.

tetrahydrozoline hcl, zinc sulfate solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-540
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**

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-540-01	1 in 1 CARTON	03/22/2023	
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	03/22/2023	

Labeler - United Natural Foods, Inc. (943556183)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(41163-540) , pack(41163-540) , label(41163-540)

Revised: 12/2023 United Natural Foods, Inc.