EQUALINE MULTI-SYMPTOM EYE DROPS- polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops
United Natural Foods, Inc.

Equaline Multi-Symptom Eye Drops 15mL (PLD)

Active ingredients

Polyethylene glycol 400 1%

Tetrahydrozoline HCl 0.05%

Zinc sulfate 0.25%

Purposes

Lubricant

Redness reliever

Astringent

Uses

- relieves dryness of the eye
- for temporary relief of discomfort and redness of the eye due to minor eye irritations
- for temporary relief of burning and irritation due to exposure to wind or sun
- for protection against further 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
- overuse may cause more eye redness
- remove contact lenses before using
- do not touch tip of container to any surface to avoid contamination
- replace cap after using
- do not use if this solution changes color or becomes cloudy

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye lasts

- condition worsens or lasts 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
- children under 6 years of age: ask a doctor

Other information

- some users may experience a brief tingling sensation
- store at 20°-25°C (68°-77°F)

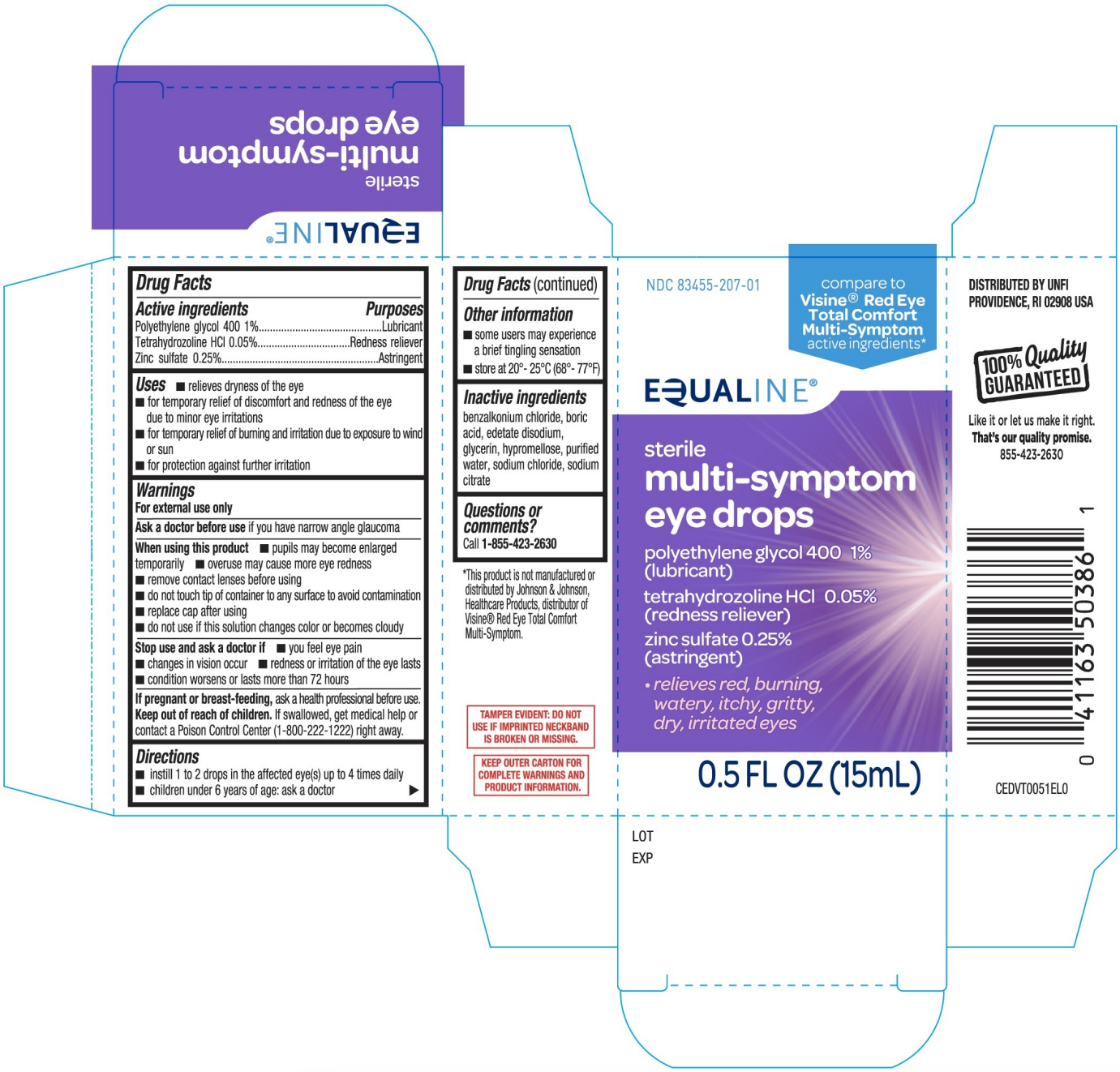
Inactive ingredients

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

Questions or comments?

Call 1-855-423-2630

Equaline Multi-Symptom Eye Drops 15mL



EQUALINE MULTI-SYMPTOM EYE DROPS

polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83455-207
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC SULFATE (UNII: 89DS0H96TB) (ZINC CATION - UNII:13S1S8SF37)	ZINC SULFATE	0.25 g in 100 mL
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ) (POLYETHYLENE	POLYETHYLENE GLYCOL	1 g

GLYCOL 400 - UNII:B697894SGQ)	400	in 100 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83455-207-01	1 in 1 BOX	06/20/2023	
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	06/20/2023	

Labeler - United Natural Foods, Inc. (943556183)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	pack(83455-207) , manufacture(83455-207) , label(83455-207)

Revised: 2/2024 United Natural Foods, Inc.