

**GOOD NEIGHBOR PHARMACY EYE DROPS MULTI-SYMPTOM RELIEF-
polyethylene glycol 400, tetrahydrozoline hcl, zinc sulfate solution/ drops
Amerisource Bergen Drug Corp.**

GNP Eye Drops Multi-Symptom Relief 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)

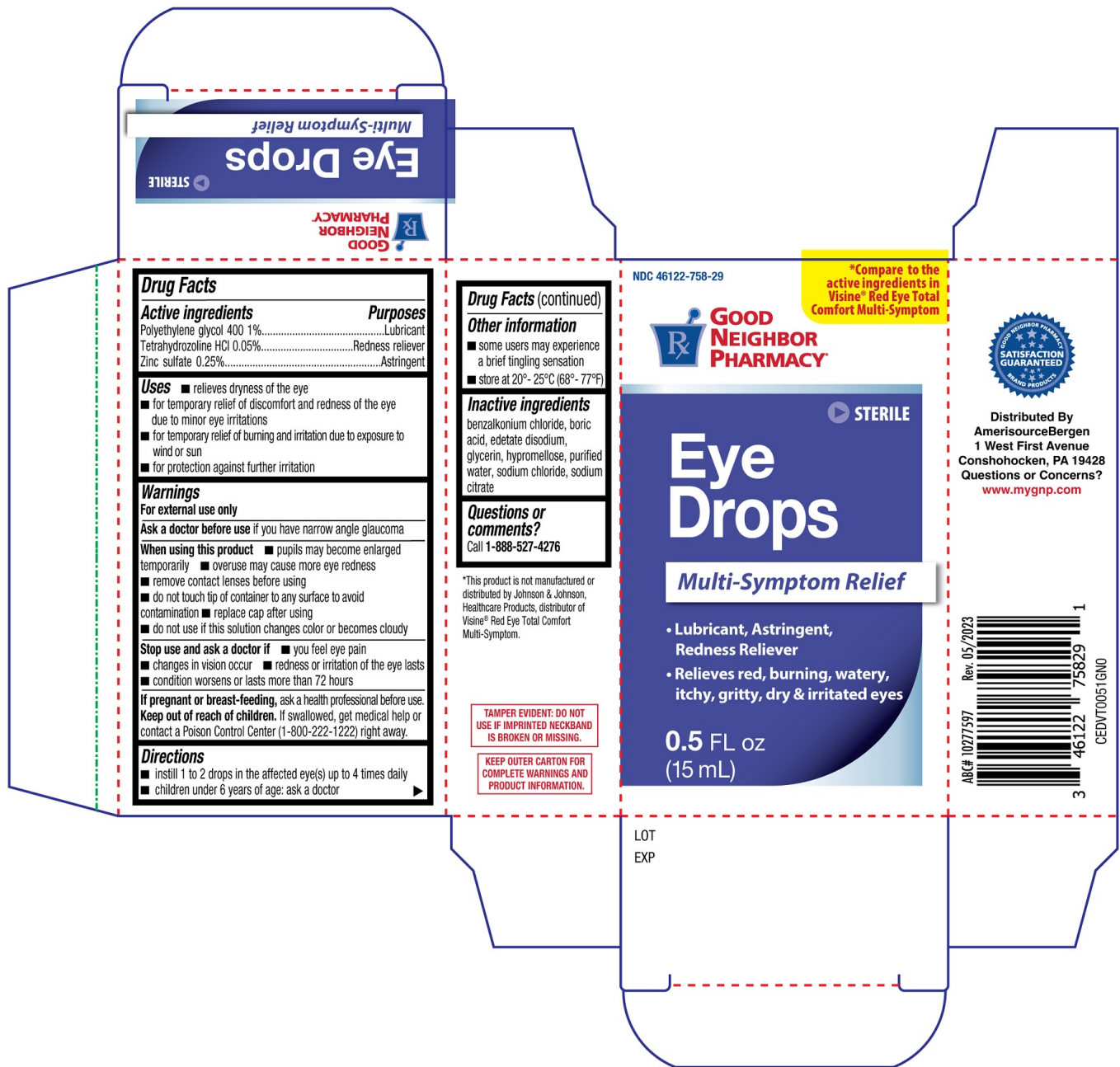
Inactive ingredients

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

Questions or comments?

Call 1-888-527-4276

GNP Eye Drops Multi-Symptom Relief 15mL



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-758
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: B6978945GQ) (POLYETHYLENE GLYCOL,	POLYETHYLENE GLYCOL	1 g

UNSPECIFIED - UNII:3WJQ0SDW1A)	400	in 100 mL
TETRAHYDROZOLINE HYDROCHLORIDE (UNII: 0YZT43HS7D) (TETRAHYDROZOLINE - UNII:S9U025Y077)	TETRAHYDROZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-758-29	1 in 1 BOX	06/05/2023	
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	06/05/2023	

Labeler - Amerisource Bergen Drug Corp. (007914906)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(46122-758) , pack(46122-758) , label(46122-758)

Revised: 12/2023

Amerisource Bergen Drug Corp.