GOODSENSE REDNESS RELIEF PLUS- gllycerin, naphazoline hydrochloride solution
Geiss, Destin & Dunn, Inc.

GoodSense Redness Relief Plus (PLD)

Active ingredients

Glycerin.....0.5%

Naphazoline hydrochloride.....0.03%

Purpose

Glycerin ....Lubricant

Naphazoline hydrochloride...Redness reliever

Uses

- for the relief of redness of the eye due to minor eye irritations
- for the temporary relief of burning and irritation due to dryness of the eye
- for use as a protectant against further irritation or dryness of the eye

Warnings

For external use only

Do not use if solution changes color or becomes cloudy

Ask a doctor before use if you have narrow angle glaucoma

When using this product

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become enlarged temporarily

Stop use and ask a doctor if

- you experience eye pain
- you experience changes in vision
- you experience continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

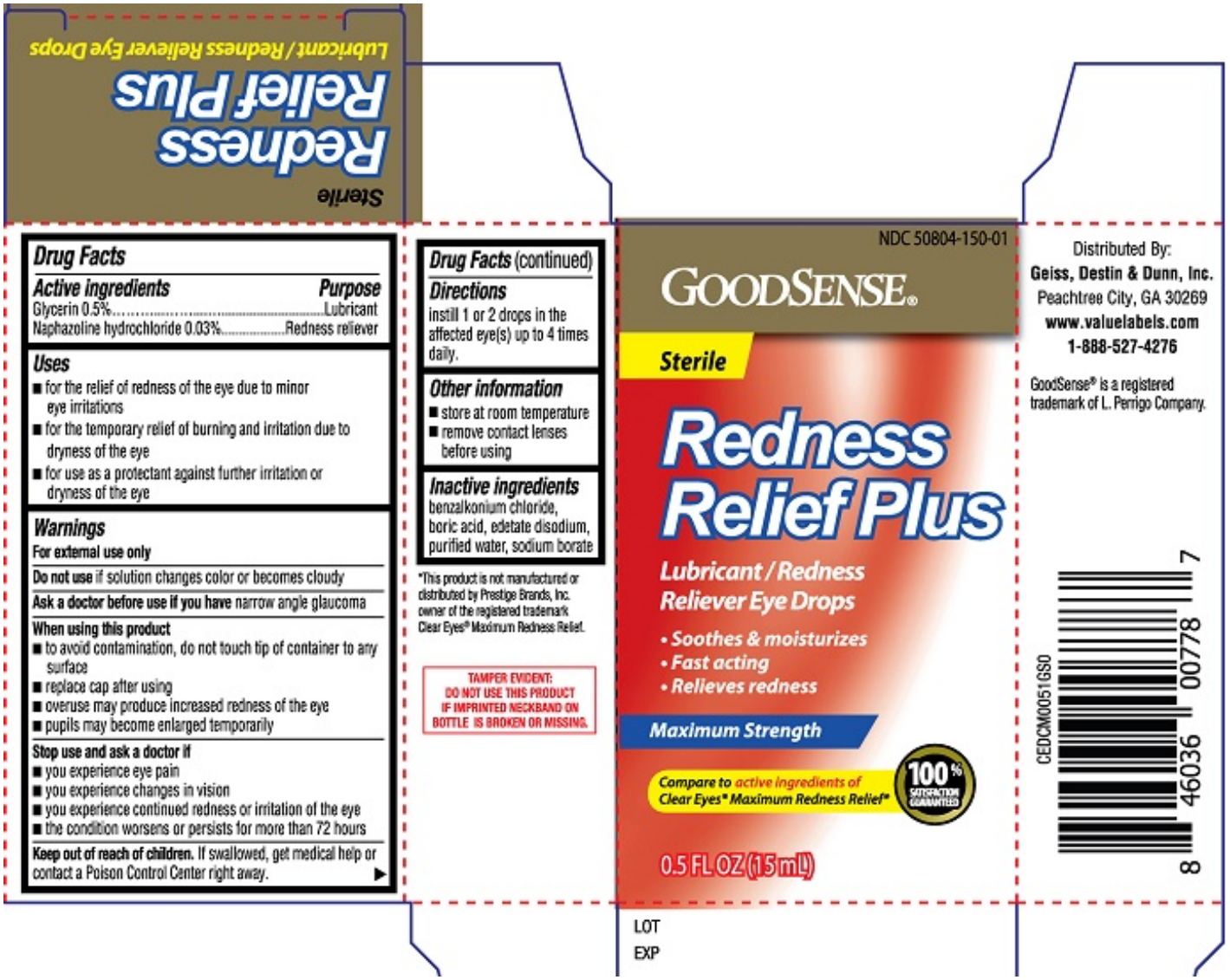
Instill 1 or 2 drops in the affected eye(s) up to 4 times daily.

Other information

- store at room temperature
- remove contact lenses before using

Inactive ingredients

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



GOODSENSE REDNESS RELIEF PLUS

gllycerin, naphazoline hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-150
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.03 g in 100 mL
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50804-150-01	1 in 1 CARTON	01/02/2020	
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	01/02/2020	

Labeler - Geiss, Destin & Dunn, Inc. (076059836)

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

Establishment

Name	Address	ID/FEI	Business Operations
K.C. Pharmaceuticals, Inc.		174450460	manufacture(50804-150) , pack(50804-150) , label(50804-150)

Revised: 12/2025 Geiss, Destin & Dunn, Inc.