

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

GoodSense Redness Relief Plus (PLD)

Active ingredients

Glycerin.....0.5%

Naphazoline hydrochloride.....0.03%

Purpose

GlycerinLubricant

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

Drug Facts

Active ingredients	Purpose
Glycerin 0.5%	Lubricant
Naphazoline hydrochloride 0.03%	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.

Drug Facts (continued)

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

*This product is not manufactured or distributed by Prestige Brands, Inc. owner of the registered trademark Clear Eyes® Maximum Redness Relief.

TAMPER EVIDENT:
DO NOT USE THIS PRODUCT IF IMPRINTED NECKBAND ON BOTTLE IS BROKEN OR MISSING.

GOODSENSE®

Sterile

Redness Relief Plus

Lubricant / Redness Reliever Eye Drops

- Soothes & moisturizes
- Fast acting
- Relieves redness

Maximum Strength

Compare to active ingredients of Clear Eyes® Maximum Redness Relief*

100% SATISFACTION GUARANTEED

0.5 FL OZ (15 mL)

NDC 50804-150-01

Distributed By:
Geiss, Destin & Dunn, Inc.
Peachtree City, GA 30269
www.valuelabels.com
1-888-527-4276

GoodSense® is a registered trademark of L. Perrigo Company.

LOT
EXP

CEDCN0051GSD

8 46036 00778 7

GOODSENSE REDNESS RELIEF PLUS

glycerin, 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: 059QF0K00R)	
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.