

**SIGNATURE CARE EYE DROPS REDNESS AND DRY EYE RELIEF- glycerin,
naphazoline hydrochloride solution/ drops
Better Living Brands LLC**

Signature Care Eye Drops Redness and Dry Eye 15 mL (PLD)

Active ingredients

Glycerin 0.25%

Naphazoline hydrochloride 0.012%

Purposes

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
- changes in vision occur
- redness or irritation of the eye lasts
- condition worsens
- symptoms last for more than 72 hours

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 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

Eye Drops Redness & Dry Eye Relief
Signature Care
Quality Guaranteed

Drug Facts
Active ingredients
Glycerin 0.25%.....Lubricant
Naphazoline hydrochloride 0.012%.....Redness reliever
Purposes

Uses
■ for the relief of redness of the eye due to minor eye irritations
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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 (1-800-222-1222) 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
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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®.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED NECKBAND IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Scan here for more information

NDC 21130-703-01

COMPARE TO the active ingredients in Clear Eyes®*

STERILE
Eye Drops Redness & Dry Eye Relief
Lubricant & Redness relief
Soothes & Protects

0.5 FL OZ (15 mL)

LOT
EXP

DISTRIBUTED BY
BETTER LIVING BRANDS LLC
P.O. BOX 99
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www.betterlivingbrandsLLC.com

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SIGNATURE CARE EYE DROPS REDNESS AND DRY EYE RELIEF

glycerin, naphazoline hydrochloride solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-703
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C00X) (GLYCERIN - UNII:PDC6A3C00X)	GLYCERIN	0.25 g in 100 mL
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.012 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-703-01	1 in 1 CARTON	10/05/2018	
1		15 mL in 1 BOTTLE, PLASTIC; 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	10/05/2018	

Labeler - Better Living Brands LLC (009137209)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(21130-703) , pack(21130-703) , label(21130-703)