CAREONE MAXIMUM REDNESS RELIEF EYE DROPS- glycerin, naphazoline hci solution/ drops
Retail Business Services, LLC

CareOne Maximum Redness Relief Eye Drops (PLD)

Active ingredients

Glycerin 0.5%

Naphazoline HCl 0.03%

Purposes

Lubricant

Redness reliever

Uses

- for the relief of redness of the eye due to minor eye irriatations
- 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 (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

Questions or comments?

Call 1-888-527-4276

CareOne Maximum Redness Relief Eye Drops 15mL

CAREONE MAXIMUM REDNESS RELIEF EYE DROPS

glycerin, naphazoline hci solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72476-012
Route of Administration	OPHTHALMIC		

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

NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.03 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72476-012-01	1 in 1 BOX	01/12/2020	
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	01/12/2020	

Labeler - Retail Business Services, LLC (967989935)

Registrant - KC Pharmaceuticals, Inc. (174450460)

Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(72476-012) , pack(72476-012) , label(72476-012)

Revised: 12/2023 Retail Business Services, LLC