MEIJER REDNESS RELIEF- glycerin, naphazoline hci solution/ drops Meijer Distribution, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Meijer Redness Relief (Glycerin, Naphazoline HCI)

Active Ingredients

Glycerin 0.25%

Naphazoline HCI 0.012%

Purposes

Lubricant

Redness Reliever

Uses

- relieves redness of the eye due to minor eye irritations
- for use as a protectant against further irritation or to relieve dryness of the eye
- for the temporary relief of burning and irritation due to 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 irriation 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 to 2 drops in the affected eye(s) up to 4 times daily

Other information

store at room temperature

Inactive ingredients

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

Questions or comments?

Call 1-888-527-4276

Meijer Redness Relief



MEIJER REDNESS RELIEF

glycerin, naphazoline hci solution/ drops

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:41250-846 Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII: H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.012 g in 100 mL	
		0.25 %	

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

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:41250-846- 01	1 in 1 BOX	04/27/2020			
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 final	part349	04/27/2020		

Labeler - Meijer Distribution, Inc. (006959555)

$\pmb{Registrant -} \ \ \mathsf{KC} \ \mathsf{Pharmaceuticals}, \ \mathsf{Inc.} \ (174450460)$

Establishment			
Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(41250-846), pack(41250-846), label(41250-846)

Revised: 4/2020 Meijer Distribution, Inc.