WALGREENS MAXIMUM REDNESS RELIEF- glycerin, naphazoline hydrochloride solution WALGREEN COMPANY

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.

Naphazoline HCL 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 the dryness of the eye.
- for the use as a protectant against further irritation or dryness of the eye.

Warnings

For external use only.

Do not use if solution changes 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

- to open bottle, push cap down and twist counter-clockwise. To close bottle, twist clockwise until it stops turning.
- put 1 to 2 drops in the affected eye(s) up to 4 times daily

Other information

- store at room temperature (15° to 30°C 59° to 86°F)
- remove contact lenses before using

Inactive ingredients

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

DISTRIBUTED BY:

WALGREEN CO.

200 WILMOT RD.
DEERFIELD, IL 60015
walgreens.com
Made in Korea



WALGREENS MAXIMUM REDNESS RELIEF

glycerin, naphazoline hydrochloride solution					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:0363-1465		363-1465	
Route of Administration	ОРНТНАЬМІС				
A . T . 1' ./A .' 36 '					
Active Ingredient/Active Moiety					
Ingredient Name		Basis of Strengt	th	Strength	
CL VCEDIN (LINIII, DDCC A2CACV) (CL)	VCEDIN LIMII.DDCC A2COOV)		CLVCEDIN		0.0025 mg

GETGERIN (ONII, PDCUASCUOA) (GETGERIN - ONII, PDCUASCUOA)	GLICERIN	in 1 mL
NAPHAZOLINE HYDRO CHLO RIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII: H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.00012 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)			
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)			
BORIC ACID (UNII: R57ZHV85D4)			
SODIUM BORATE (UNII: 91MBZ8H3QO)			
WATER (UNII: 059QF0KO0R)			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0363-1465-15	1 in 1 CARTON			
1		15 mL in 1 BOTTLE, DROPPER			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part349	11/25/2014		

Labeler - WALGREEN COMPANY (008965063)

Revised: 11/2014 WALGREEN COMPANY