EYE WASH- water solution CMC Group 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.

Eye Wash

Drug Facts

Active ingredient

Purified Water 99.1%

Purpose

Eyewash

\Box Use

• For cleansing the eye to help relieve irritation or burning by removing loose foreign material

□ Warnings

IFor external use only.

Do not use

if solution changes color or becomes cloudy

When using this product

- to avoid contamination, do not touch tip of container to any surface
- do not reuse
- once opened, discard
- obtain immediate medical treatment for all open wounds in or near the eyes

Stop use and ask a doctor if

you experience:

- eye pain
- changes in vision
- continued redness
- irritation of the eye
- condition worsens or perisists

Keep out of reach of children.

If swallowed, get medical help of contact a Poison Control Center right away.

Directions

• Flush the affected eyes as needed, controlling the rate of flow of solution by pressure on the bottle

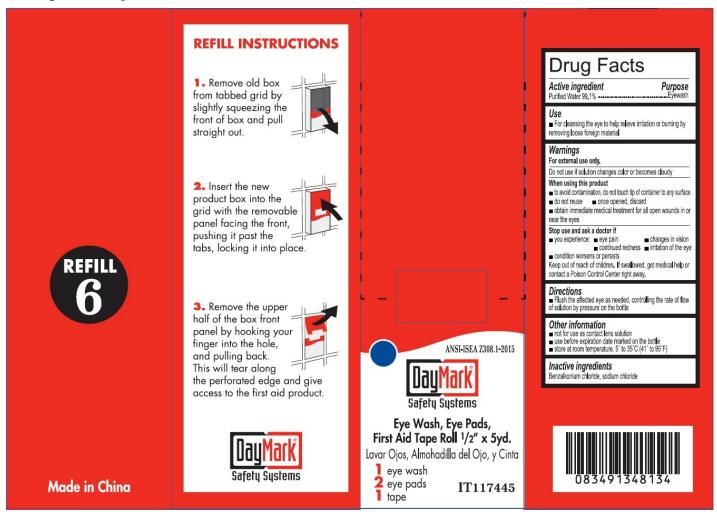
Other information

- not for use as contact len solution
- use before expiration date marked on the bottle
- store at room temperature, 5 loll to 35 loll (41 loll to 95 loll)

Inactive ingredients

Benzalkonium chloride, sodium chloride

Package Labeling:



EYE WASH water solution **Product Information** HUMAN OTC DRUG NDC:49687-0010 Product Type Item Code (Source) **Route of Administration OPHTHALMIC Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER $991\,mg$ in $1\,mL$

Inactive Ingredients				
Ingredient Name	Strength			
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:49687-0010-1	1 in 1 BOX	08/09/2016				
1		30 mL in 1 TUBE; 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	08/09/2016			

Labeler - CMC Group Inc. (117201448)

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