PHYSICIANSCARE EYE WASH- water solution Acme United Corporation

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.

Physicians Care Eye Wash

Active Ingredient

Purified Water 98%

Purpose

Eyewash

Indications:

For flushing the eyes to help relieve irritation, discomfort, burning, stinging, or itching by removing

loose foreign material and air pollutants (smog or pollen).

Warnings: For External Use Only

- To avoid contamination, do not touch tip of bottle to any surface
- Do not reuse
- Discard bottle after use

Do Not Use:

- If solution changes color or becomes cloudy
- With contact lenses
- If bottle is open or seal is broken

Stop use and consult a doctor if you experience:

- Eye pain
- Changes in vision
- Continued redness or irritation of the eye or if the condition worsens or persists

Obtain immediate medical treatment for all open wounds in or near the eyes .

Keep out of reach of children.

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

Directions:

- To pour hold bottle securely, twist off top to remove
- Control rate of flow by pressure on the bottle
- Flush the affected eye(s) as needed
- Do not touch bottle tip to eye
- If necessary, continue flushing with emergency eyewash or shower.

Inactive Ingredients:

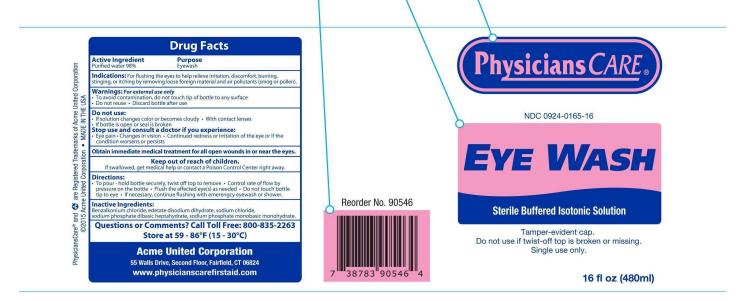
Benzalkonium chloride, edetate disodium dihydrate, sodium chloride, sodium phosphate dibasic

heptahydrate, sodium phosphate monobasic monohydrate.

Questions or Comments? Call Toll Free: 800-835-2263

Store at 59 - 86°F (15 - 30°C)

All PINK areas will print white Ink



PHYSICIANSCARE EYE WASH water solution **Product Information Product** Type HUMAN OTC DRUG NDC:0924-0165(NDC:57349-913) Item Code (Source) **OPHTHALMIC Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R) WATER 0.98 mL in 1 mL **Inactive Ingredients Ingredient** Name Strength BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) EDETATE DISODIUM (UNII: 7FLD91C86K) SODIUM CHLORIDE (UNII: 451W47IQ8X) SODIUM PHO SPHATE, DIBASIC, HEPTAHYDRATE (UNII: 70 WT22SF4B) SO DIUM PHO SPHATE, MO NO BASIC, MO NO HYDRATE (UNII: 593YOG76 RN) Packaging Item Code **Package Description** Marketing Start Date Marketing End Date # 1 NDC:0924-0165-16 480 mL in 1 BOTTLE; Type 0: Not a Combination Product 12/05/2014 2 NDC:0924-0165-32 946 mL in 1 BOTTLE; Type 0: Not a Combination Product 12/05/2014 **3** NDC:0924-0165-01 30 mL in 1 BOTTLE; Type 0: Not a Combination Product 12/05/2014

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part349	12/05/2014				
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Labeler - Acme United Corporation (001180207)

Establishment						
Name	Address	ID/FEI	Business Operations			
Acme United Corporation		045924339	relabel(0924-0165), repack(0924-0165)			

Establishment						
Name	Address	ID/FEI	Business Operations			
Acme United Corporation		080119599	relabel(0924-0165), repack(0924-0165)			

Revised: 1/2016

Acme United Corporation