EYEWASH STATION ADDITIVE CONCENTRATE- chlorhexidine gluconate and propylene glycol liquid
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.

PhysiciansCare Eyewash Station Additive Concentrate

Product Information

Ingredients

Chlorhexidine gluconate 1% w/v Purified Water, Propylene Glycol

Purpose

Preservative

Use

A preservative for use in potable self-contained emergency eyewash stations

Warnings

For external use only. In case of contact with eye in undiluted form, flush with clean water

Do not use
- in full strength

When using this product
- do not change dilution or use with other chemicals
- do not reuse

Stop use and ask a doctor if *you have*
- changes in vision
- eye pain
- condition worsens or persists
- continued redness or irritation of the eye

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions
- wear protective eyewear and gloves

- clean potable eyewash station and rinse with potable water
- partially fill station with potable water
- remove tamper evident seal and cap from bottle
- add entire contents of the bottle to the eyewash station container
- fill the station to the manufacturer's required level
- date and initial inspection tag
- station should be cleaned and refilled every 120 days when using this product
- in advance of emergency, add the concentrate to potable water to have a solution available

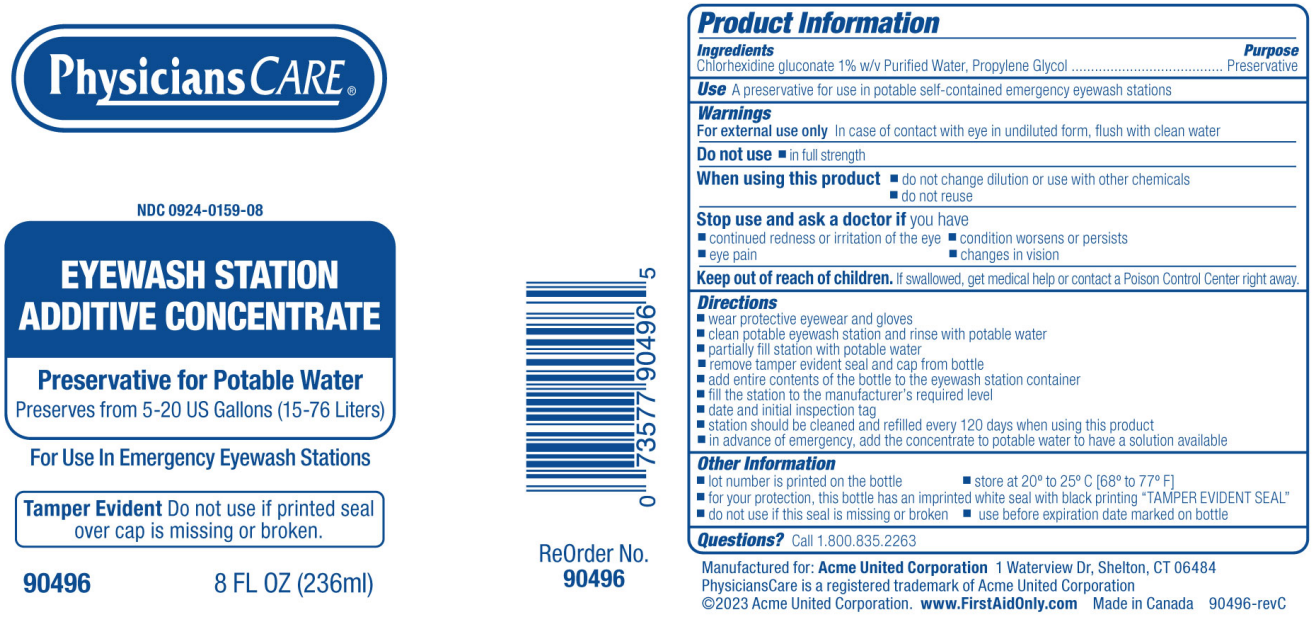
Other information

- lot number is printed on the bottle
- store at 20° to 25° C [68° to 77° F]
- for your protection, this bottle has an imprinted white seal with black printing "TAMPER EVIDENT SEAL"
- do not use if this seal is missing or broken
- use before expiration date marked on bottle

Questions ?

☎ Call 1.800.835.2263

Bottle Label



EYEWASH STATION ADDITIVE CONCENTRATE

chlorhexidine gluconate and propylene glycol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-0159(NDC:65785-035)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E) (CHLORHEXIDINE - UNII:R4KO0DY52L)	CHLORHEXIDINE GLUCONATE	145.6 kg in 2800 L
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	280 kg in 2800 L

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	2374.4 L in 2800 L

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0159-08	0.236 L in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	01/09/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part349	01/09/2023	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0159) , repack(0924-0159)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	repack(0924-0159) , relabel(0924-0159)

Revised: 1/2023 Acme United Corporation