PHYSICIANSCARE OPHTHALMIC SOLUTION EYEWASH- purified water 98.3% solution
Acme United Corporation

PhysiciansCare Ophthalmic Solution Eyewash

Active Ingredients

Purified Water 98.3%

Purpose

Eyewash

Use

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

Warnings

For external use only

Do not use
- if you experience any open wounds in or near the eyes and obtain immediate medical treatment
- 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

Stop use and ask a doctor if you have any of the following
- 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
- remove tamper evident seal and cap
- flush the affected eye as needed, controlling the rate of flow of solution by pressure on the bottle.

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

Inactive ingredients

boric acid, sodium borate, sodium chloride

Questions? Call 1-800-835-2263

Physicians Care 8oz NDC 0924-0162 New Label 11-2018

PHYSICIANSCARE OPHTHALMIC SOLUTION EYEWASH

purified water 98.3% solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-0162(NDC:65785-162)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	929 g in 946 mL

Inactive Ingredients

Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0162-08	236 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	06/28/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022305	06/24/2013	

Labeler - Acme United Corporation (001180207)

Registrant - Niagara Pharmaceuticals, Inc. (205477792)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0162) , repack(0924-0162)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation Vancouver Division		080119599	relabel(0924-0162) , repack(0924-0162)

Establishment

Name	Address	ID/FEI	Business Operations
Niagara Pharmaceuticals, Inc.		205477792	manufacture(0924-0162)

Revised: 11/2019 Acme United Corporation