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

Physicians Care 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

	CADE	Drug Facts	
(Physicia	ns CARE 。	Active ingredient 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	
NDC 0924	-0162-08	Do not use if you experience any open wounds in or near the eyes and obtain imn if solution changes color or becomes cloudy	nediate medical treatment
		When using this product = to avoid contamination, do not touch tip of co do not reuse once opened, discard	ntainer to any surface
EYEV	VASH	Stop use and ask a doctor if you have any of the following • continued redness or irritation of the eye • condition worsens or persists • eye pain • changes in vision	
Desifie d Ma	lan 00 00/	Keep out of reach of children. If swallowed, get medical help or contact a right away.	Poison Control Center
Purified Wa Ophthalmic Sol		Directions remove tamper evident seal and cap flush the affected eye a controlling the rate of flow of solution by pressure on the bottle	s needed,
Single Use	Sterile Solution	Other Information I to number is printed on the bottle store at 20° to for your protection, this bottle has an imprinted white seal with black printing "TAM do not use if this seal is missing or broken use before expiration date marked	IPER EVIDENT SEAL"
Temper Fuident De r	act use if printed appl	Inactive ingredients boric acid, sodium borate, sodium chloride	
Tamper Evident Do r over cap is miss		Questions? Call 1.800.835.2263	
24-050	8 FL OZ (236ml)	ReOrder No. Manufactured for: Acme United Corporation 55 Walls Dr, Fairfield, CT O 24-050 PhysiciansCare is a registered trademark of Acme United Corporation ©2018 Acme United Corporation. www.FirstAidOnly.com	

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

PHYSICIANSCARE OF	HTHALMIC SO	LUTION EY	EWA	SH		
purified water 98.3% solution						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:0924-		NDC:0924-0	0162(NDC:65785-162)	
Route of Administration	OPHTHALMIC					
Active Ingredient/Active M	niety					
	dient Name		Basis	of Strength	n Strength	
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)WATER			0	929 g	g in 946 mL	
The stine Transit diamon						
Inactive Ingredients	In and is no Norma				C	4
BORIC ACID (UNII: R57ZHV85D4)	Ingredient Name				Strength	
SODIUM BORATE (UNII: 91MBZ8H3QO)						
SODIUM CHLORIDE (UNII: 451W47IQ8X)						
Packaging						
# Item Code	Package Descriptio	n	Ma	Marketing Start Date		arketing End Date
1 NDC:0924-0162- 08 236 mL in 1 BC Product	TTLE, UNIT-DOSE; Type 0	: Not a Combination	n 06/28	06/28/2013		

Marketing Information						
Marketing Category	ting Category Application Number or Monograph Citation		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