PUR-WASH- water solution Niagara Pharmaceuticals Inc.

Pur-Wash

Drug Facts

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

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

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

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
- replace with sterile eyecup provided
- avoid contamination of rim and inside surfaces of the eyecup
- place eyecup surface to the affected eye, pressing tightly to prevent the escape of the liquid and tilt the head backward
- open eyelids wide and rotate eyeball while controlling the rate of flow of solution by pressure on the bottle to ensure thorough bathing with the wash

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 905 690-6277 9 a.m. to 5 p.m. EST Mon-Fri

Manufactured by: Niagara Pharmaceuticals Inc. 60 Innovation Dr. Flamborough ON L9H 7P3

PRINCIPAL DISPLAY PANEL - 946 mL Bottle Label

Pur-Wash

Purified Water, 98.3% Ophthalmic Solution Eyewash

Single Use

Manufactured by: Niagara Pharmaceuticals Inc. 60 Innovation Dr. Flamborough ON L9H 7P3 Made in Canada

Sterile Solution

32 Fl Oz [946 mL]

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water solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:65785-164

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 CHLORIDE (UNII: 451W47IQ8X) SODIUM BORATE (UNII: 91MBZ8H3QO)

Item Code Package Description Marketing Start Date Marketing End Date 1 NDC:65785-164- 01 Product 09/12/2011 2 NDC:65785-164- 02 Product 09/12/2011

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
NDA	NDA022305	09/12/2011			

Labeler - Niagara Pharmaceuticals Inc. (205477792)

Establishment								
Name	Address	ID/FEI	Business Operations					
Niagara Pharmaceuticals Inc.		205477792	manufacture(65785-164)					

Revised: 10/2018 Niagara Pharmaceuticals Inc.