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

• pull cover off cap

• 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]
- do not use if tamper evident ring is 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 With
Sterile Eyecup Attached

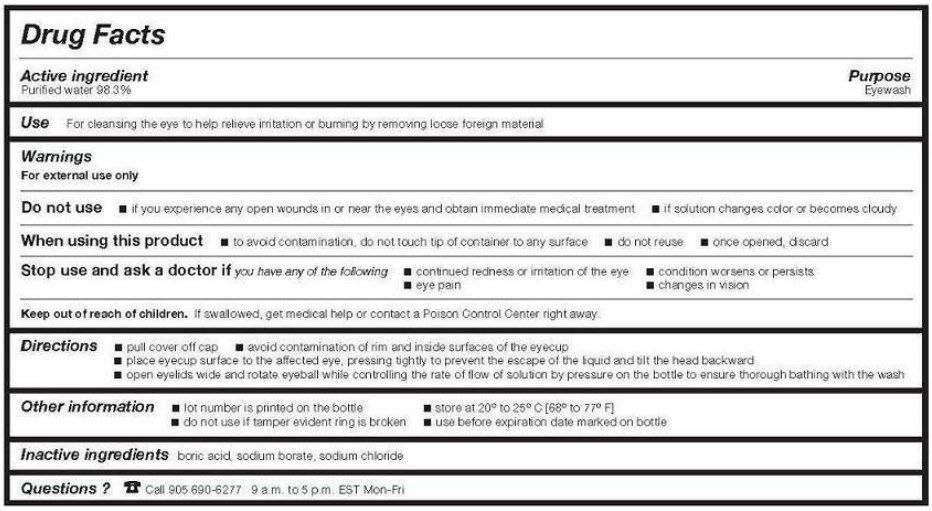
NDC 65785-166-02

Single Use

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

Sterile Solution

32 Fl Oz [946 mL]



PUR-WASH

water solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65785-166
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)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65785-166-01	473 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	05/24/2013	
2	NDC:65785-166-02	946 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	05/24/2013	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
NDA	NDA022305	05/24/2013	

Labeler - Niagara Pharmaceuticals Inc. (205477792)

Establishment

Name	Address	ID/FEI	Business Operations
Niagara Pharmaceuticals Inc.		205477792	manufacture(65785-166)

Revised: 10/2018 Niagara Pharmaceuticals Inc.