FICHA TÉCNICA

PUR-WASH™
AGUA PURIFICADA
98,3%

Propósito: Lavado de ojos, solución oftálmica, lavado ocular. Uso único
Solución estéril
Tamaño: 16 fl oz (473mL) / 32 fl oz (946 mL)
Ingrediente Activo: Agua purificada 98,3%
Ingredientes Inactivos: Ácido bórico, borato de sodio, cloruro de sodio.

NDC 65785-168-02
Fabricado por: Niagara Pharmaceuticals Inc.
60 Innovation Dr., Flamborough ON L9H 7P3
Fabricado en Canadá

Advertencias: Solo para uso externo
-No usar
- Si experimenta alguna herida abierta en o cerca de los ojos y recibe tratamiento médico inmediato.
- Si la solución cambia de color o se vuelve turbia

-Al usar este producto
- Para evitar la contaminación, no toque la punta del recipiente con ninguna superficie
- no reutilizar
- una vez abierto, deseche

Uso: Para limpiar el ojo para ayudar a aliviar la irritación o el ardor al eliminar el material extraño suelto.
- El número de lote está impreso en la botella
- Almacenar de 20° a 25° C [68° a 77° F]
- Para su protección, esta botella tiene un sello blanco impreso con la impresión negra "TAMPER EVIDENT SEAL"
- No usar si falta este sello o está roto
- Usar antes de la fecha de caducidad marcada en la botella

-Deje de usarlo y pregunte a un médico si tiene alguno de los siguientes:
- Enrojecimiento o irritación continua del ojo
- La condición empeora o persiste
- Dolor de ojos
- Cambios en la visión

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6982-7308 / 386-4386 ventas1@safetysolutionsglobal.com LLano Bonito, Plaza Vista marina, Local #2, 3, 4.

PUR-WASH- eyewash solution
Niagara Pharmaceuticals, Inc.

Niagara Pur-Wash

Active ingredient

Purified water 98.3%

Purpose

Eyewash

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

Use

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

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
- avoid contamination of rim and inside surfaces of the nozzle

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- place nozzle to the affected area and tilt the bottle to pour liquid out through the nozzle
- 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 1-800-615-2216 9 a.m. to 5 p.m. EST Mon-Fri

Dosage and Administration

- remove tamper evident seal and cap
- avoid contamination of rim and inside surfaces of the nozzle
- place nozzle to the affected area and tilt the bottle to pour liquid out through the nozzle
- 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

Warnings

For external use only

Package Label.Principal Display Panel

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PUR-WASH

eyewash solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65785-168
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	465 mL in 473 mL

Inactive Ingredients

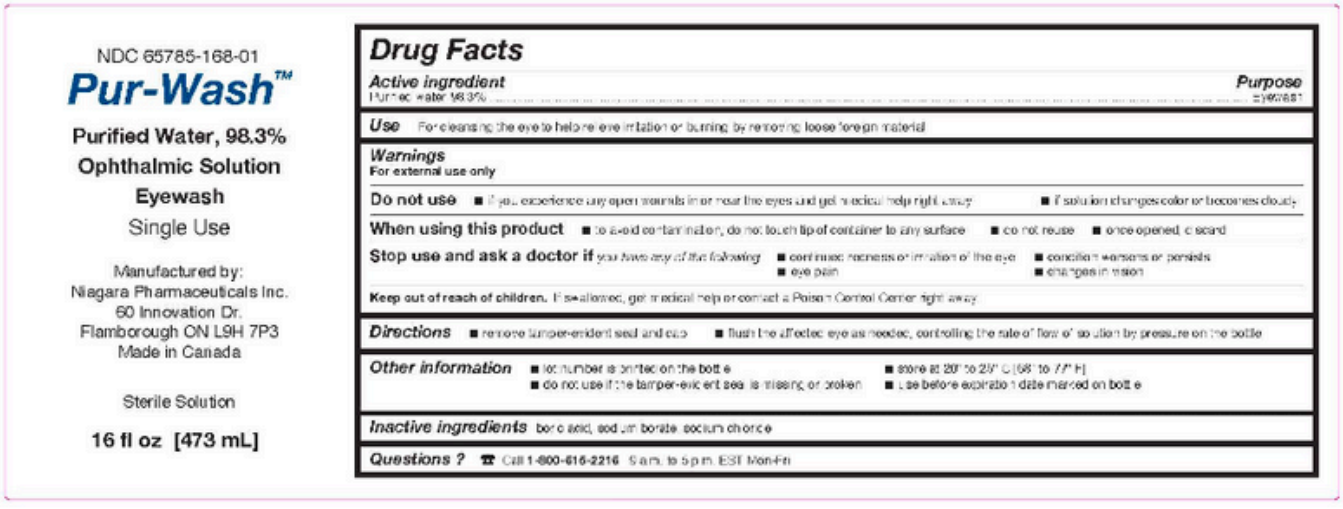
Ingredient Name	Strength
BORIC ACID (UNII: R57ZHV85D4)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65785-168-01	473 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	09/01/2015	
2	NDC:65785-168-02	946 mL in 1 BOTTLE, UNIT-DOSE; Type 0: Not a Combination Product	09/01/2015	



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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022305	01/30/2015	

Labeler - Niagara Pharmaceuticals, Inc. (205477792)

Registrant - Niagara Pharmaceuticals, Inc. (205477792)

Establishment

Name	Address	ID/FEI	Business Operations
Niagara Pharmaceuticals, Inc.		205477792	manufacture(65785-168)

Revised: 7/2025 Niagara Pharmaceuticals, Inc.

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