EYEWASH- water solution
MWI

APEXA EYEWASH

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 experience*

- 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

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 888-628-0581

Principal Display Panel Text for Container Label:

NDC 13985-609-04

Ophthalmic

Solution

Eyewash

Purified Water, 98.3%

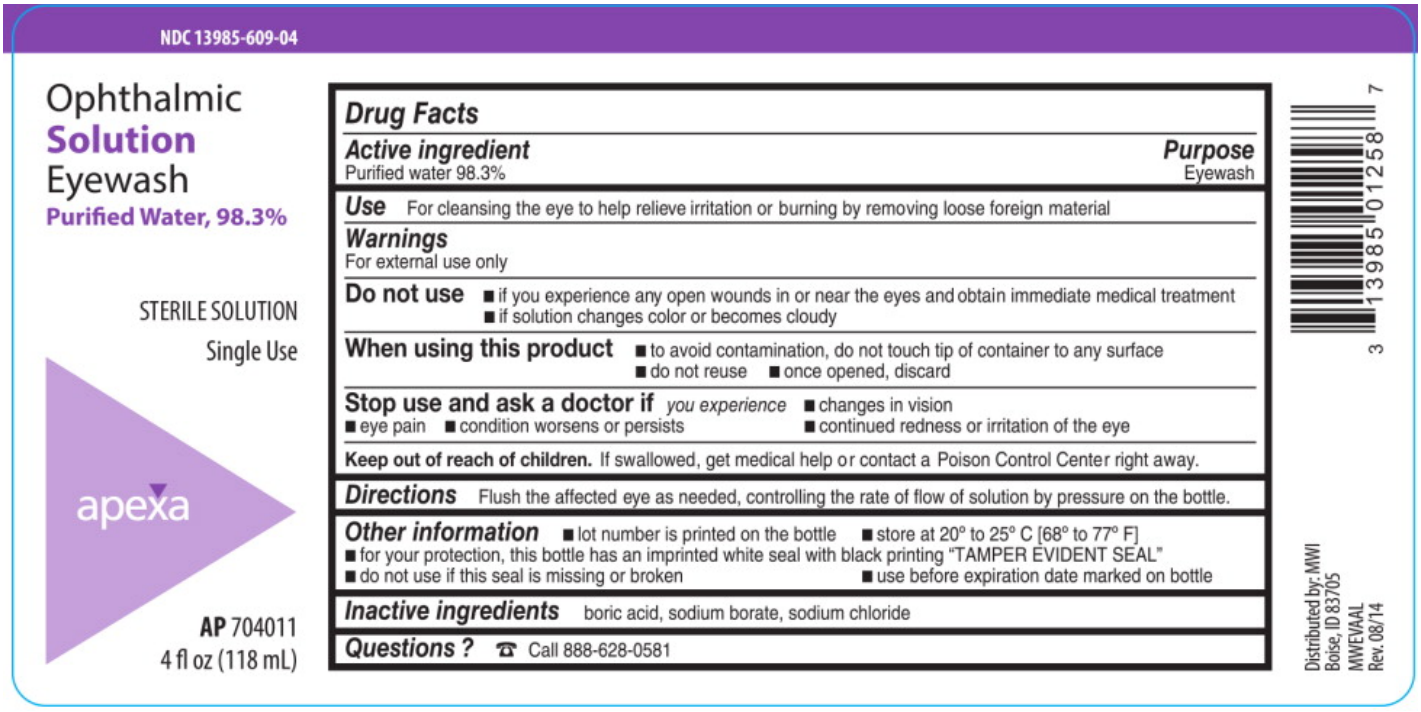
STERILE SOLUTION

Single Use

APEXA logo

AP 704011

4 fl.oz. (118 mL)



EYEWASH

water solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13985-609
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:13985-609-04	1 in 1 BOTTLE	03/25/2015	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022305	02/23/2015	

Labeler - MWI (019926120)

Registrant - Akorn Operating Company LLC (117693100)

Establishment

Name	Address	ID/FEI	Business Operations
Niagara Pharmaceuticals, Inc.		205477792	manufacture(13985-609)

Revised: 10/2020 MWI