

EYEWASH- water solution

Akorn

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click [here](#).

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:

Akorn logo

OPHTHALMIC

SOLUTION

EYEWASH

Purified Water, 98.3%

NDC 59399-001-04

Single Use

FOR ANIMAL USE ONLY

Distributed by:

Akorn Operating Company LLC

Gurnee, IL

60031

EVAAL Rev. 07/22

Made in

Canada

Sterile

Solution

4 fl.oz. (118 mL)



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EYEWASH

water solution

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:59399-001
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:59399-001-04	1 in 1 BOTTLE		
1		118 mL in 1 BOTTLE		
2	NDC:59399-001-08	1 in 1 BOTTLE		
2		236 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		02/01/2015	

Labeler - Akorn (117693100)

Revised: 7/2022

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