

**EYE WASH- water solution**  
**GFA Production (Xiamen) Co., Ltd.**

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**Eye Wash**

**Drug Facts**

***Active ingredient***

Purified Water 99.1%

**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 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 • obtain immediate medical treatment for all open wounds in or near the eyes

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

**Stop use and ask a doctor if**

• you experience: • eye pain • changes in vision • continued redness • irritation of the eye • condition worsens or persists

***Directions***

• Flush the affected eye as needed, controlling the rate of flow of solution by pressure on the bottle

## Other information

- not for use as contact lens solution
- use before expiration date marked on the bottle
- store at room temperature, 5° to 35°C (41° to 95°F)

## Inactive ingredients

Benzalkonium chloride, sodium chloride

## Package Labeling:



**REFILL  
6**

### REFILL INSTRUCTIONS

- 1.** Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.


- 2.** Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.


- 3.** Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.





ANSI-ISEA Z308.1-2015



**Eye Wash, Eye Pads,  
First Aid Tape Roll 1/2" x 5yd.**  
Lavar Ojos, Almohadilla del Ojo, y Cinta

**1** eye wash  
**2** eye pads  
**1** tape

**IT117445**

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## EYE WASH

water solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50814-010
<b>Route of Administration</b>	OPHTHALMIC		

<b>Active Ingredient/Active Moiety</b>				
<b>Ingredient Name</b>		<b>Basis of Strength</b>	<b>Strength</b>	
<b>WATER</b> (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)		WATER	991 mg in 1 mL	
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)				
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50814-010-01	1 in 1 BOX	08/09/2016	12/31/2025
1		30 mL in 1 TUBE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M003	08/09/2016	12/31/2025	

**Labeler** - GFA Production (Xiamen) Co., Ltd. (421256261)

Revised: 10/2024

GFA Production (Xiamen) Co., Ltd.