EYEWASH- purified water liquid
Reliance Medical Ltd

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Eyewash

Drug Facts

Active ingredient

Purified Water 99.1 %

Purpose

Eyewash

Uses

For cleansing the eye to remove loose foreign material.

Warnings

For external use only.

Do not use

• if solution changes colour or becomes cloudy

• for injection

• as a contact lens solution.

When using the product

• do not re-use, once opened, discard.

• to avoid contamination, do not touch the tip of the container to any surface.

Stop use and consult a doctor if

• you experience eye pain.

• changes in vision.

• continued redness or irritation of the eye.

• the condition worsens or persists.

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 Poison Control Centre right away.

Directions

• smoothly twist the lid in either direction to break the seal and remove the lid. • position the bottle above the eye and gently squeeze.

• flush the affected eye as needed controlling the rate of flow of solution by pressure on the bottle. • cleanse the eye(s) with as much solution as required.

Other information

• store between 5°C and 35°C (41° to 95° F).

• use before expiry date.

• do not use if seal is broken prior to use.

Inactive ingredient

Sodium chloride.

Questions?

Call 1-844-405-7233. Monday - Friday, 9am - 5pm. You may also report serious side effects to this number.

MANUFACTURED FOR:

Reliance Medical LTD, Radnor Park Trading

Estate, Congleton, Cheshire, England, CW12 4XP, UK.

IMPORTED AND DISTRIBUTED BY:

Innovative Safety Supply USA Inc. 5150 West 76th St. Indianapolis,

IN, 46268 USA.

MADE IN CHINA

Compliance:

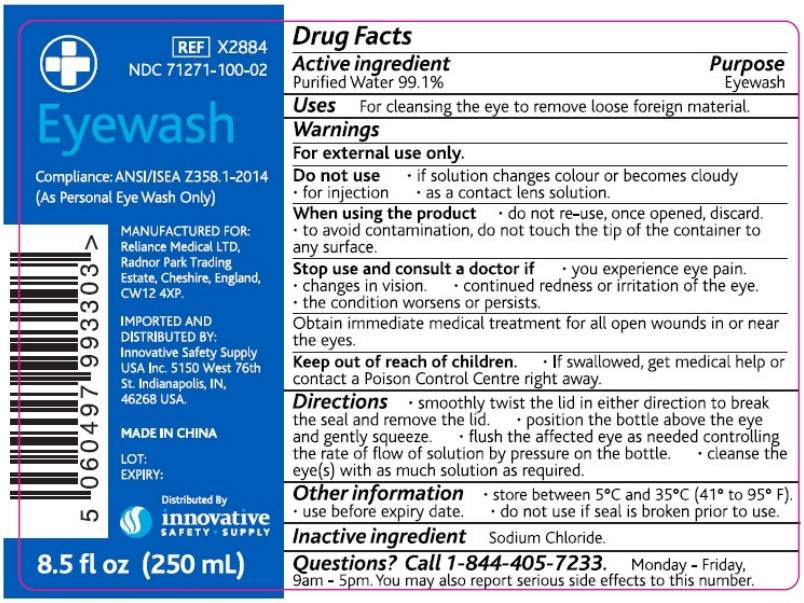
ANSI/ISEA Z358.1-2014

(As Personal Eye Wash Only)

Packaging

20 mL

250 mL



EYEWASH

purified water liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71271-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	0.991 mL in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71271-100-01	25 in 1 CARTON	07/03/2017	
1		20 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:71271-100-02	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/03/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	07/03/2017	

Labeler - Reliance Medical Ltd (349044854)

Revised: 1/2018 Reliance Medical Ltd