A-MED BRAND FIRST AID EYE AND SKIN-RINSE- purified water solution
Oliver Landon Intl Inc.

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.

A-Med Wash First Aid Eye Rinse

Drug Facts

Active ingredient

Purified water USP 98.577% w/v

Purpose

Eyewash

Use for flushing the eye to remove loose foreign material.

Warnings

For external use only

Do not use
- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes. Get medical help right away.

When using this product
- do not touch tip of container to any surface to avoid contamination
- do not reuse; once opened, discard

Stop use and ask a doctor if
- 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.

Inactive ingredients disodium disodium EDTA, polyhexanide 0.0001% (preservative), sodium chloride, sodium phosphate dibasic dodecahydrate, sodium phosphate, monobasic dihydrate. Sodium hydroxide may be used to adjust pH.

Distributed by: Oliver Landon Intl Inc.

21 Pine Road, Belleville, St. Michael

BB 11113 Barbados

Questions or concerns:

email - info@oliverlandon.biz or call (800) 839-5929

Made in U.S.A.

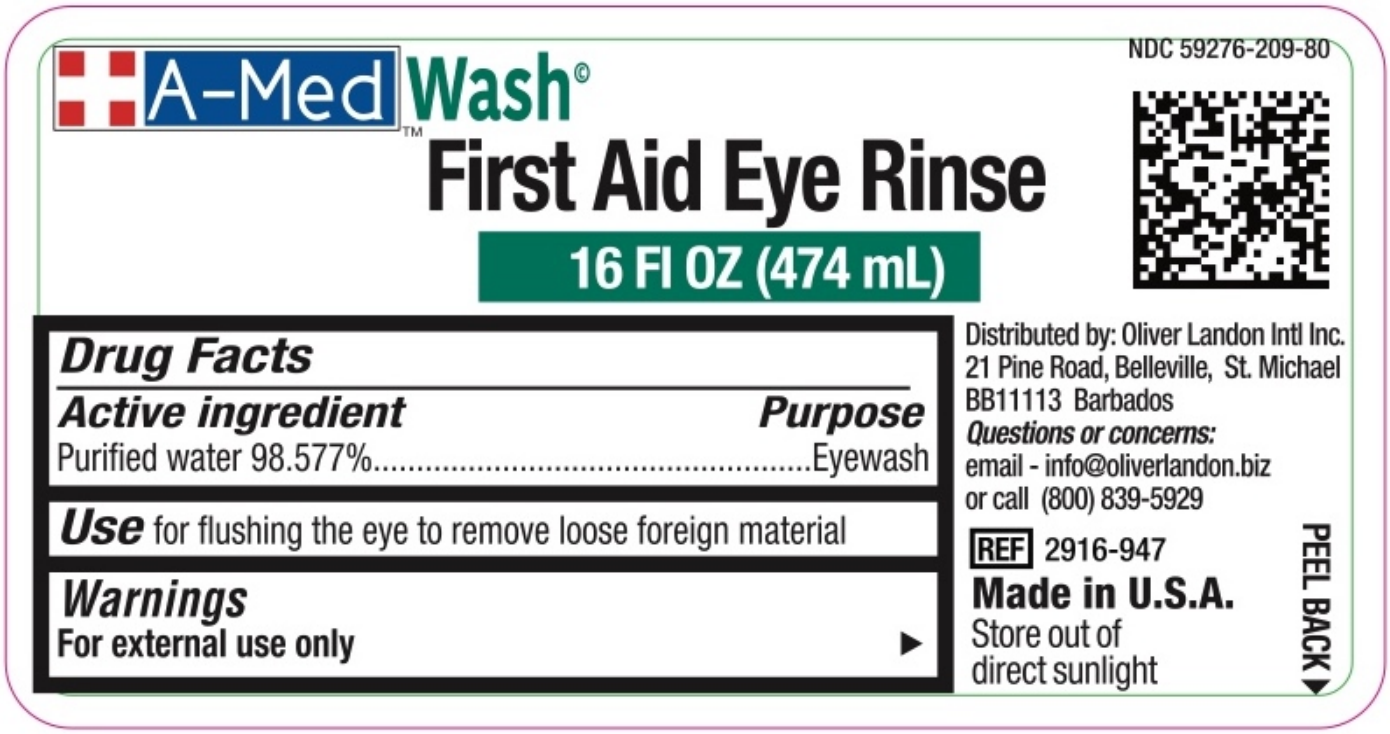
Store out of direct sunlight.

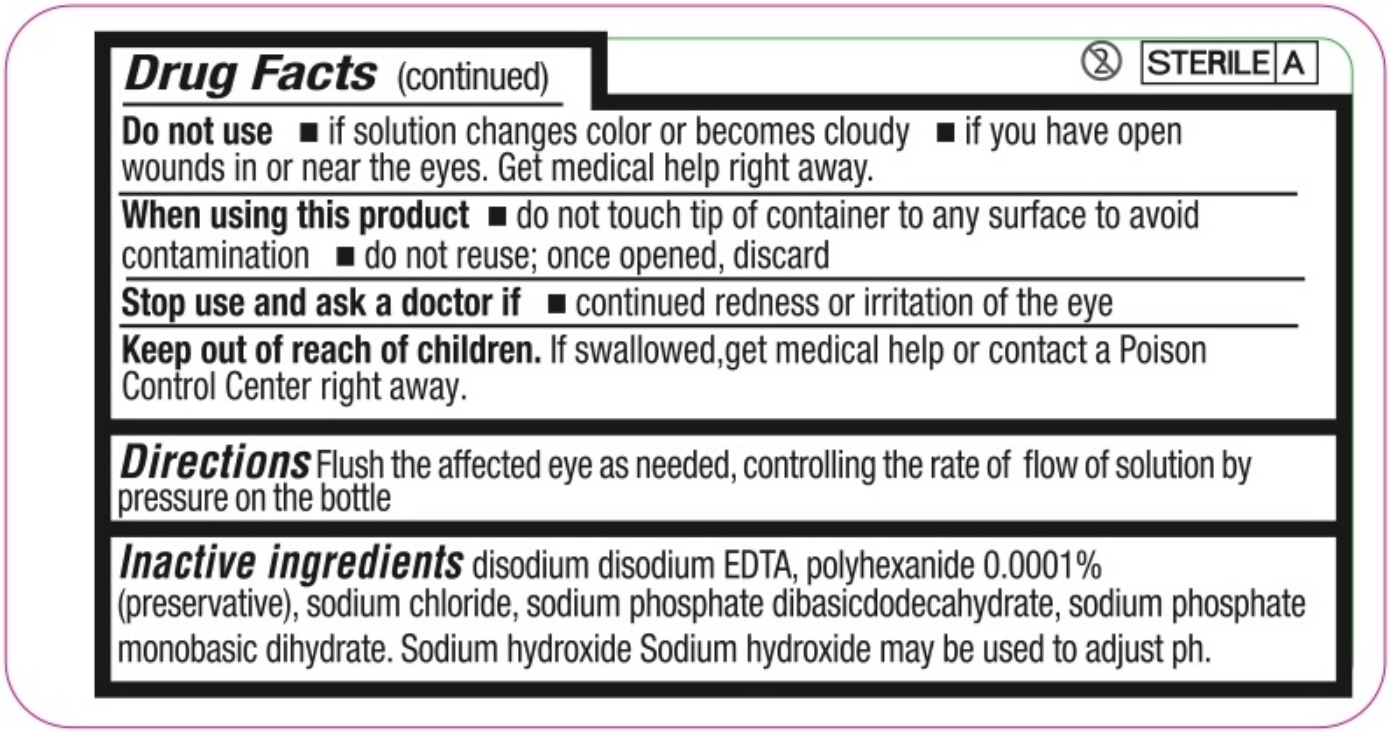
A-Med Wash

NDC 59276-209-80

First Aid Eye Rinse

16 FL OZ (474 mL)





A-MED BRAND FIRST AID EYE AND SKIN-RINSE

purified water solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59276-209
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.577 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
POLIHEXANIDE (UNII: 322U039GMF)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM PHOSPHATE, DIBASIC, DODECAHYDRATE (UNII: E1W4N241FO)	
SODIUM PHOSPHATE, MONOBASIC, DIHYDRATE (UNII: 5QWK665956)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59276-209-12	5 in 1 BOX	11/01/2013	
1		15 mL in 1 VIAL; Type 0: Not a Combination Product		
2	NDC:59276-209-14	10 in 1 BOX	11/01/2013	12/20/2020
2		15 mL in 1 VIAL; Type 0: Not a Combination Product		
3	NDC:59276-209-16	30 in 1 BOX	11/01/2013	12/21/2020
3		15 mL in 1 VIAL; Type 0: Not a Combination Product		
4	NDC:59276-209-18	60 in 1 BOX	11/01/2013	12/20/2020
4		15 mL in 1 VIAL; Type 0: Not a Combination Product		
5	NDC:59276-209-21	4 in 1 BOX	11/01/2013	12/20/2020
5		30 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
6	NDC:59276-209-31	1 in 1 BOX	11/01/2013	12/20/2020
6		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
7	NDC:59276-209-41	1 in 1 BOX	11/01/2013	12/20/2020
7		118 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
8	NDC:59276-209-50	148 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2020	
9	NDC:59276-209-61	1 in 1 BOX	11/01/2013	12/20/2020
9		236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

10	NDC:59276-209-71	1 in 1 BOX	11/01/2013	12/20/2020
10		350 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product		
11	NDC:59276-209-81	1 in 1 BOX	11/01/2013	12/20/2020
11		474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
12	NDC:59276-209-82	2 in 1 BOX	11/01/2013	12/20/2020
12		474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
13	NDC:59276-209-91	1 in 1 BOX	11/01/2013	12/20/2020
13		947 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
14	NDC:59276-209-92	2 in 1 BOX	11/01/2013	12/20/2020
14		947 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
15	NDC:59276-209-22	4 in 1 BOX	12/20/2020	
15		15 mL in 1 VIAL; Type 0: Not a Combination Product		
16	NDC:59276-209-30	30 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product	12/20/2020	
17	NDC:59276-209-40	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2020	
18	NDC:59276-209-60	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2020	
19	NDC:59276-209-70	350 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	12/20/2020	
20	NDC:59276-209-90	947 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2020	
21	NDC:59276-209-80	474 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2020	
22	NDC:59276-209-10	15 mL in 1 VIAL; Type 0: Not a Combination Product	11/01/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	11/01/2013	

Labeler - Oliver Landon Intl Inc. (815240195)

Revised: 12/2021 Oliver Landon Intl Inc.