WHITE RAIN ANTIBACTERIAL HAND SPRING WATER- benzalkonium chloride soap
International Wholesale, Inc.

White Rain Antibacterial Hand Soap, Spring Water

Drug Facts:

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

for hand washing, reduces germs on the skin

Warnings

for external use only-hands only

Do Not use

on damaged or broken skin.

When using the product

keep out of eyes. Rinse with water to remove.

Stop and consult a doctor if

irritation and redness develops.

Keep out of reach of children.

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

Directions:

Wash Hands and rinse.

Inactive Ingredients

Water, Cetrimonium Chloride, Glycerin, Lauramidopropylamine Oxide and Myristamidopropylamine Oxide, Sodium Chloride, Cocamide MEA, PEG-120 Methyl

Glucose Dioleate, Citric Acid, Fragance, Disodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Red 33, Blue 1

Other Information

Avoid Storing at extreme temperatures

Package Labeling:

WHITE RAIN ANTIBACTERIAL HAND SPRING WATER

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52862-626
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	

LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52862-626-00	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/01/2025	

Labeler - International Wholesale, Inc. (080442566)

Registrant - International Wholesale, Inc. (080442566)

Revised: 5/2025 International Wholesale, Inc.