

WHITE RAIN ANTIBACTERIAL HAND LAVENDER BREEZE- benzalkonium chloride liquid
International Wholesale, Inc.

White Rain Antibacterial Hand Soap LAVENDER BREEZE

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

for hand washing to decrease bacteria on the skin. ***External***

Warnings

for external use only.

When using this product

avoid contact with eyes. In case of eye contact, flush with water.

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:

Pump into DRY hands, Lather vigorously for at least 15 seconds. Rinse and dry thoroughly.


Inactive Ingredients

Inactive Matter: Water(Aqua), Lauramine Oxide, Coco-Glucoside, Glyceryl Oleate, Cetrimonium Chloride, PEG-150 Distearate, Myristamine Oxide, Glycerin, Fragrance(Parfum), Citric Acid, Sodium Chloride, Disodium EDTA, Benzophenone-4, Methylchloroisothiazolinone, Methylisothiazolinone, FD&C Red No. 33, FD&C Blue No. 1.

Package Labeling:



White Rain



Antibacterial Hand Soap

Mild and gentle antibacterial soap for everyday cleanliness

Lavender Breeze
mild & gentle


7.5 fl oz (221ml)

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DISTRIBUTED BY: **INNOVATIVE BRANDS** 4000 ALLEN RD
ALLEN PARK MI 48101
1-888-346-6688

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Represents transparency

WHITE RAIN ANTIBACTERIAL HAND LAVENDER BREEZE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52862-622
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
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SULISOBENZONE (UNII: 1W6L629B4K)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
WATER (UNII: 059QF0KO0R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52862-622-00	1 in 1 CASE	02/01/2023	
1		221 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/01/2023	

Labeler - International Wholesale, Inc. (161872676)

Revised: 11/2023

International Wholesale, Inc.