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

White Rain Foaming Antibacterial Hand Soap Spring Water

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

External

for hand washing to decrease bacteria on the skin.

Warnings

for external use only.

When using this product

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

Stop use 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:

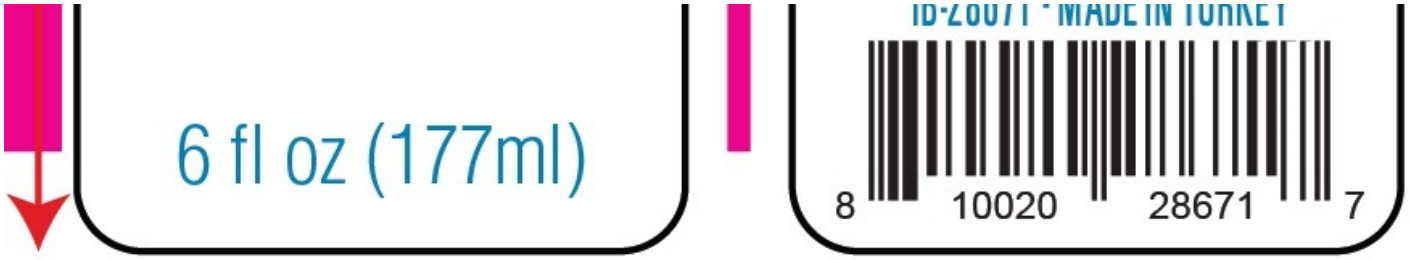
Pump into DRY hands. Lather vigorously for at least 15 seconds. Rinse and dry thoroughly.

Inactive Ingredients

Water(Aqua), Lauramine Oxide, Cetrimonium Chloride, Myristamine Oxide, Glycerin, Coco-Glucoside, Glyceryl Oleate, Fragrance(Parfum), Citric Acid, Sodium Chloride, Disodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, FD&C Blue No.1,

FD&C Red 33.

Package Labeling:



Represents transparency

WHITE RAIN FOAMING ANTIBACTERIAL HAND SPRING WATER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52862-615
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)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLYCERYL OLEATE (UNII: 4PC054V79P)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52862-615-00	1 in 1 CASE	01/01/2023	
1		177 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)	01/01/2023	

Labeler - International Wholesale, Inc. (161872676)

Revised: 11/2023 International Wholesale, Inc.