

**WHITE RAIN ANTIBACTERIAL HAND WHITE TEA- benzalkonium chloride liquid
International Wholesale, Inc.**

White Rain Antibacterial Hand Soap WHITE TEA

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

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, Methylchloroisoithiazolinone, Methylisoithiazolinone, FD&C Yellow No. 4, FD&C Blue No. 1.

Package Labeling:



| | |
|---|--|
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DISTRIBUTED BY: **INNOVATIVE BRANDS** 4000 ALLEN RD
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Represents transparency

WHITE RAIN ANTIBACTERIAL HAND WHITE TEA

benzalkonium chloride liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:52862-623 |
| 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 |
|---|----------|
| WATER (UNII: 059QF0KO0R) | |
| 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) | |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) | |

Packaging

| # | Item Code | Package Description | Marketing Start | Marketing End |
|---|-----------|---------------------|-----------------|---------------|
|---|-----------|---------------------|-----------------|---------------|

| # | Item Code | Package Description | Date | Date |
|---|------------------|---|------------|------|
| 1 | NDC:52862-623-00 | 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.