# FRESH FEELS HAND SANITIZER MIST CHERRY SCENT- ethyl alcohol spray Pearl World Inc.

-----

#### FRESH FEELS HAND SANITIZER MIST CHERRY SCENT

### **Drug Facts**

### **Active Ingredient**

ETHYL ALCOHOL 70%

## **Purpose**

**ANTISEPTIC** 

#### Uses

- To decrease bacteria on the skin that could cause disease.
- When water, soap and towel are not available.
- Recommended for repeated use.

# Warnings

For external use only. Flammable. Keep away from fire or flame.

# When using this product:

• Keep out of eyes, ears, or mouth. • In case of eye contact, flush eyes thoroughly with water. • Avoid contact with broken skin. • Do not inhale or ingest.

**Stop use and ask a doctor if •** Redness or irritation develop. • Condition persists for more than 72 hours.

# Keep out of reach of children.

- Children should be supervised by an adult when using this product.
- If swallowed, get medical help or contact a Poison Control Center right away.

### **Directions**

- Spray product in your palm and rub hands together until dry.
- For children under 6, use only under adult supervision.
- Not recommended for infants.

#### Other Information

- Do not store above 105°F. May discolor some fabrics.
- Harmful to wood finishes & plastics.

## **Inactive Ingredients**

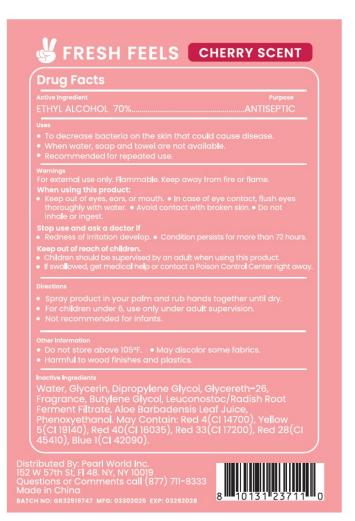
Water, Glycerin, Dipropylene Glycol, Glycereth-26, Fragrance, Butylene Glycol, Leuconostoc/Radish Root Ferment Filtrate, Aloe Barbadensis Leaf Juice, Phenoxyethanol. May Contain: Red 4(Cl 14700), Yellow 5(Cl 19140), Red 40(Cl 16035), Red 33(Cl 17200), Red 28(Cl 45410), Blue 1(Cl 42090)

Distributed by: Pearl World Inc. 152 W 57th St, Fl 48. NY, NY 10019 Made in China

Questions or Comments call (877) 711-8333

# **Packaging**





# FRESH FEELS HAND SANITIZER MIST CHERRY SCENT

ethyl alcohol spray

| Product Information     |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:69933-253 |
| Route of Administration | TOPICAL        |                    |               |

| Active Ingredient/Active Moiety                        |                          |                 |
|--|--------------------------|-----------------|
| Ingredient Name  | <b>Basis of Strength</b> | Strength        |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL                  | 70 mL in 100 mL |

| Inactive Ingredients  |          |  |
|---|----------|--|
| Ingredient Name   | Strength |  |
| WATER (UNII: 059QF0KO0R)                                    |          |  |
| GLYCERIN (UNII: PDC6A3C0OX)                                 |          |  |
| DIPROPYLENE GLYCOL (UNII: E107L85C40)                       |          |  |
| GLYCERETH-26 (UNII: NNE56F2N14)                             |          |  |
| BUTYLENE GLYCOL (UNII: 3XUS85K0RA)                          |          |  |
| LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458) |          |  |
| ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)                     |          |  |
| PHENOXYETHANOL (UNII: HIE492ZZ3T)                           |          |  |
| FD&C RED NO. 4 (UNII: X3W0AM1JLX)                           |          |  |
| FD&C YELLOW NO. 5 (UNII: I753WB2F1M)                        |          |  |
| FD&C RED NO. 40 (UNII: WZB9127XOA)                          |          |  |
| D&C RED NO. 33 (UNII: 9DBA0SBB0L)                           |          |  |
| <b>D&amp;C RED NO. 28</b> (UNII: 767IP0Y5NH)                |          |  |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)                          |          |  |

| F | Packaging            |   |                         |                       |  |  |
|---|----------------------|---|-------------------------|-----------------------|--|--|
| # | Item Code            | Package Description   | Marketing Start<br>Date | Marketing End<br>Date |  |  |
| 1 | NDC:69933-<br>253-12 | 1 in 1 PACKET   | 04/17/2025              |                       |  |  |
| 1 |                      | 40 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product |                         |                       |  |  |

| Marketing Information |   |                         |                       |  |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |  |
| OTC Monograph Drug    | 505G(a)(3)                                  | 04/17/2025              |                       |  |
|                       |   |                         |                       |  |

# Labeler - Pearl World Inc. (043130142)

Revised: 4/2025 Pearl World Inc.