

FRESH FEELS HAND SANITIZER MIST SEASHELL SUNSHINE SCENT- ethyl alcohol spray
Amber Glow Cosmetics Limited

FRESH FEELS HAND SANITIZER MIST SEASHELL SUNSHINE 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 and 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 40 Lake (CI 16035), Red 33 Lake (CI 17200), Red 4 Lake (CI 14700), Yellow 5 Lake (CI 19140:1), Blue 1 Lake (CI 42090:2).

Fresh Feels hand sanitizer mist is lightweight, leaving hands feeling clean, soft and fragranced.

Set Contains:
HAND SANITIZER 40 ML/1.35 FL.OZ

Distributed by:
Amber Glow Cosmetics Limited
Unit 1806, 18/F.,
9 Wing Hong Street,
Cheung Sha Wan, Hong Kong
Questions or Comments call (877) 711-8333
Made in China

PHthalates FREE

GLUTEN FREE

PARABEN FREE

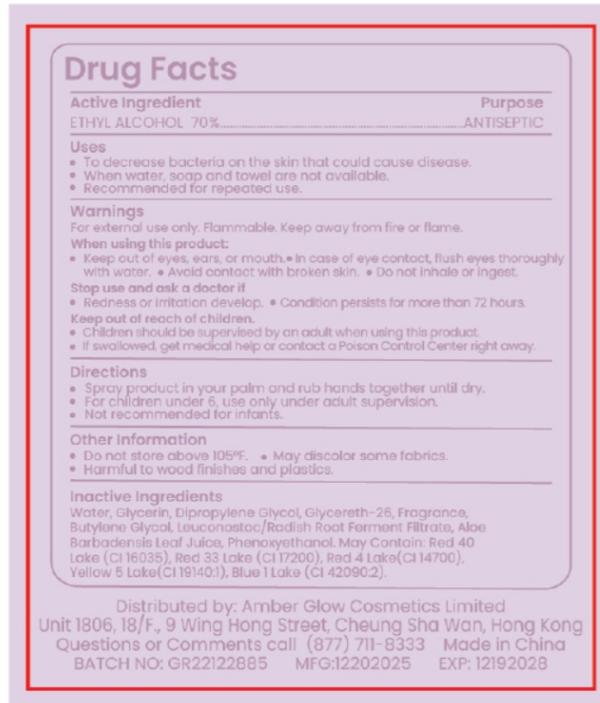
Packaging

Outer Package Label





Inner Package Label



FRESH FEELS HAND SANITIZER MIST SEASHELL SUNSHINE SCENT

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87302-202
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	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: 3XUS85KORA)	
LEUCONOSTOC/RADISH ROOT FERMENT FILTRATE (UNII: D2QHA03458)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 ALUMINUM LAKE (UNII: JQ6BLH9FR7)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87302-202-02	1 in 1 POUCH	01/07/2026	
1		40 mL in 1 BOTTLE, SPRAY; 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)	01/07/2026	

Labeler - Amber Glow Cosmetics Limited (765280476)

Revised: 12/2025

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