FRESH FEELS HAND SANITIZER MIST VANILLA BREEZE SCENT- ethyl alcohol spray
Amber Glow Cosmetics Limited

FRESH FEELS HAND SANITIZER MIST VANILLA BREEZE 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

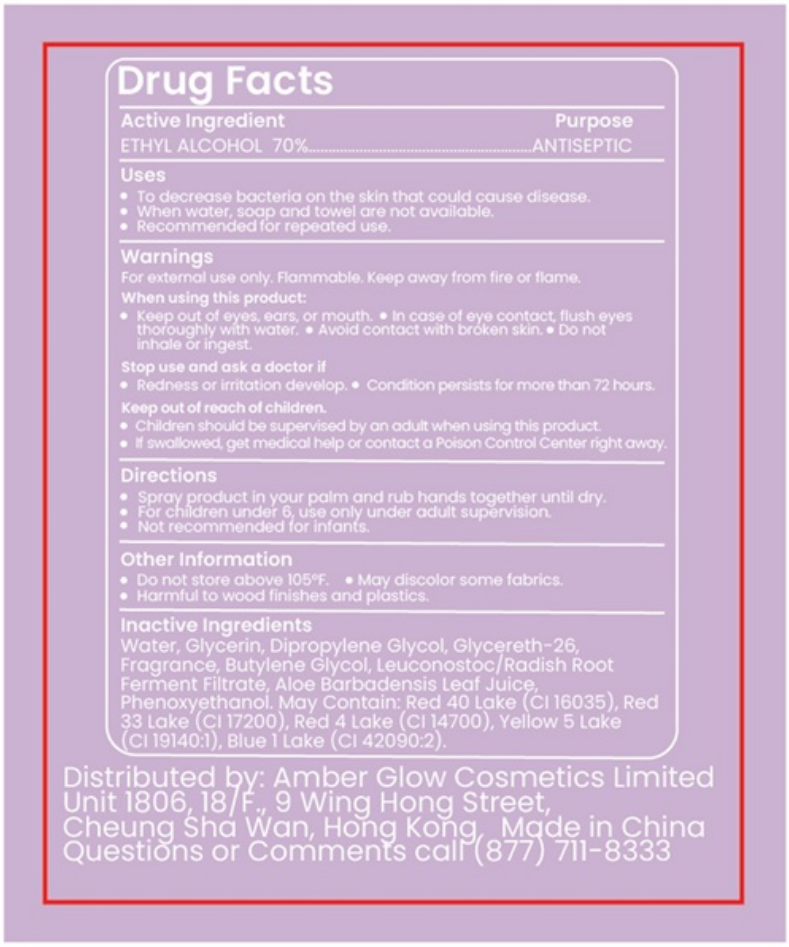
• Do not store above 105°F. • May discolor some fabrics.
• Harmful to wood finishes and plastics.

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).

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

Packaging



FRESH FEELS HAND SANITIZER MIST VANILLA BREEZE SCENT

ethyl alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87302-208

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: 3XUS85K0RA)	
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:87302-208-08	35 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/07/2026	

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 Amber Glow Cosmetics Limited