

SMART CARE VANILLA HEAVEN HAND SANITIZER- ethyl alcohol spray
Ashtel Studios, Inc.

Smart Care® VANILLA HEAVEN HAND SANITIZER

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antiseptic

Uses

Decreases bacteria on skin.

Warnings

For external use only

Flammable, keep away from fire or flame.

Do not use

- If you are allergic to any of the ingredients.

When using this product • Do not get into eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor • If irritation or rash develops and continues for more than 72 hours.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

• Adults and children 6 years and older

Spray onto hands and rub them together until dry. Recommended for repeated use.

• Children under 6 years of age

Should be supervised when using this product.

Other information

- Store between 15-30°C (59-86°F)

- Avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Gel, Aminomethyl Propanol, Blue 1, Denatonium Benzoate, Fragrance, Glycerin, PEG-40 Hydrogenated Castor Oil, Red 33, Tocopheryl Acetate, Water, Yellow 5

Questions or comments?

Call toll free **1-877-274-8358**

SCENTED & POCKET-FRIENDLY

KILLS UP TO 99% OF GERMS*

A delightful blend of creamy vanilla and sweet sugar. Indulge your senses and whisk away stress with each delicate spritz of our invigorating hand sanitizer.

Every airy mist ensures clean and nourished skin, providing a refreshing experience with each use.

*Effective at killing 99.9% *Escherichia coli*, *Staphylococcus aureus* under laboratory settings

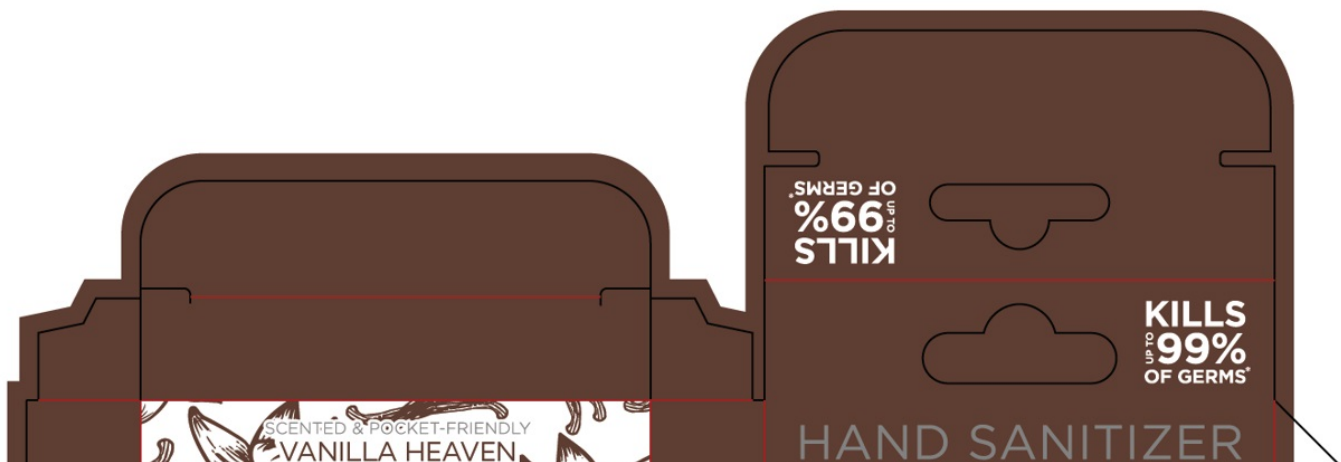
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**DISTRIBUTED BY:
ASHTEL STUDIOS, INC.
ONTARIO, CALIFORNIA 91761**

MADE IN CHINA

Packaging







SMART CARE VANILLA HEAVEN HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70108-135
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
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-135-01	1 in 1 BOX	05/05/2025	
1		30 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)	05/05/2025	

Labeler - Ashtel Studios, Inc. (148689180)

Revised: 4/2025

Ashtel Studios, Inc.