

SMART CARE SWEET AND STILL HAND SANITIZER- ethyl alcohol spray
Ashtel Studios, Inc.

Smart Care® SWEET & STILL 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, call Poison Help at 1-800-222-1222 or go to PoisonHelp.org.

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

Questions or comments?

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

KILLS UP TO 99.9% OF GERMS*

**SCENT INSPIRED BY
BURBERRY GODDESS***

SCENTED & POCKET-FRIENDLY

*SMART CARE DOES NOT USE DESIGNER BRAND FRAGRANCES AND IS NOT ASSOCIATED IN ANY WAY WITH THE DESIGNER BRANDS OR THEIR MANUFACTURERS. ALL TRADEMARKS ARE PROPERTY OF THEIR RESPECTIVE OWNER. DESIGNER/BRAND NAMES ARE SOLELY USED FOR COMPARISON PURPOSES TO GIVE CUSTOMERS AN IDEA OF FRAGRANCE CHARACTER AND SCENT ACCORDS. WE DO NOT PRESENT OUR FRAGRANCES TO BE EXACT COPIES.

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

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ASHTEL STUDIOS, INC.
ONTARIO, CALIFORNIA 91761**

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MADE IN CHINA

Packaging

Outer Package Label





Inner Package Label





SMART CARE SWEET AND STILL HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70108-192
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC 70108			

1	NDC: 70108-192-01	1 in 1 POUCH	11/01/2025	
1		50 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)	11/01/2025	

Labeler - Ashtel Studios, Inc. (148689180)