

SMART CARE SQUISHMALLOWS SWEET PEA HAND SANITIZER- ethyl alcohol spray
Ashtel Studios, Inc.

Smart Care® SQUISHMALLOWS SWEET PEA HAND SANITIZER SPRAY

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, Denatonium Benzoate, Fragrance, Glycerin, Tocopheryl Acetate, Water

Questions or comments?

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

ORIGINAL

KILLS 99.9% OF GERMS*

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

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SmartCare®

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

Packaging



Drug Facts Label

Drug Facts

Active ingredient **Purpose**
Ethyl Alcohol 70% v/vAntiseptic

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

Item # 88372-A

8 10147 88372 1

Peel ▶

Drug Facts (Continued)

Uses Decreases bacteria on skin.

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Peel ▶

Drug Facts (Continued)

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SMART CARE SQUISHMALLOWS SWEET PEA HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70108-182
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)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-182-01	28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/10/2025	

Labeler - Ashtel Studios, Inc. (148689180)

Revised: 6/2025

Ashtel Studios, Inc.