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

Smart Care[®] 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 together until dry. Recommended for repeat 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 4, Red 33, Tocopheryl Acetate, Water

Questions or comments?

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

SCENTED & POCKET-FRIENDLY

KILLS UP TO 99.9% OF MOST COMMON GERMS*

Step into spring with fresh, delicate notes of orange blossom, rose, and hyacinth. 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% of *Escherichia coli, Staphylococcus aureus* under laboratory settings

Smart Care[®] SMARTCAREUS.COM 1-909-434-0911 INTERNATIONAL

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DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

MADE IN CHINA

Packaging





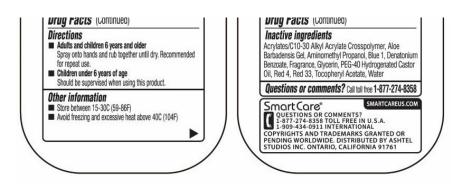


NDC: 70108-120-01



Drug Facts Label





SMART CARE SWEET PEA HAND SANITIZER

ethyl alcohol spray

Product Information					
Product Type	HUMAN OTC DRUG	AN OTC DRUG Item Code (Source) NE		NDC:70108-120	
Route of Administration	TOPICAL				
Active Ingredient/Active	e Moiety				
Ingredient Name			Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	70 mL in 100 mL	
Inactive Ingredients Ingredient Name					
	Ingredient Nam	е		Strength	
ACRYLATES/C10-30 ALKYL ACI	RYLATE CROSSPOLYMER (50000 M	PA.S) (UNII: 8Z5ZAL5H3V	()	
		50000 M	PA.S) (UNII: 8Z5ZAL5H3V	()	
ALOE VERA LEAF (UNII: ZY81Z8	3H0X)	50000 M	PA.S) (UNII: 8Z5ZAL5H3V)	
ALOE VERA LEAF (UNII: ZY81Z8 Aminomethylpropanol (UNI	3H0X) I: LU49E6626Q)	50000 M	PA.S) (UNII: 8Z5ZAL5H3V)	
ALOE VERA LEAF (UNII: ZY81Z8. AMINOMETHYLPROPANOL (UNI FD&C BLUE NO. 1 (UNII: H3R47)	3H0X) I: LU49E6626Q) <3TBD)	50000 M	PA.S) (UNII: 8Z5ZAL5H3V		
ACRYLATES/C10-30 ALKYL ACH ALOE VERA LEAF (UNII: ZY81Z8: AMINOMETHYLPROPANOL (UNII FD&C BLUE NO. 1 (UNII: H3R47) DENATONIUM BENZOATE (UNII: GLYCERIN (UNII: PDC6A3C0OX)	3H0X) I: LU49E6626Q) <3TBD)	50000 M	PA.S) (UNII: 8Z5ZAL5H3V		
ALOE VERA LEAF (UNII: ZY81Z8 AMINOMETHYLPROPANOL (UNI FD&C BLUE NO. 1 (UNII: H3R47) DENATONIUM BENZOATE (UNII:	3H0X) I: LU49E6626Q) (3TBD) : 4YK5Z54AT2)		PA.S) (UNII: 8Z5ZAL5H3V		
ALOE VERA LEAF (UNII: ZY81Z8 AMINOMETHYLPROPANOL (UNI FD&C BLUE NO. 1 (UNII: H3R47) DENATONIUM BENZOATE (UNII: GLYCERIN (UNII: PDC6A3C0OX)	3H0X) I: LU49E6626Q) <3TBD) : 4YK5Z54AT2) CASTOR OIL (UNII: 7YC686		PA.S) (UNII: 8Z5ZAL5H3V		
ALOE VERA LEAF (UNII: ZY81Z8 AMINOMETHYLPROPANOL (UNII FD&C BLUE NO. 1 (UNII: H3R47) DENATONIUM BENZOATE (UNII: GLYCERIN (UNII: PDC6A3C0OX) POLYOXYL 40 HYDROGENATED FD&C RED NO. 4 (UNII: X3W0AM	3H0X) I: LU49E6626Q) (3TBD) : 4YK5Z54AT2) D CASTOR OIL (UNII: 7YC686 1JLX)		PA.S) (UNII: 8Z5ZAL5H3V) 	
ALOE VERA LEAF (UNII: ZY81Z8 AMINOMETHYLPROPANOL (UNI FD&C BLUE NO. 1 (UNII: H3R47 DENATONIUM BENZOATE (UNII: GLYCERIN (UNII: PDC6A3C0OX) POLYOXYL 40 HYDROGENATED	3H0X) I: LU49E6626Q) <3TBD) : 4YK5Z54AT2) D CASTOR OIL (UNII: 7YC686 1JLX) BB0L)		PA.S) (UNII: 8Z5ZAL5H3V) 	

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108- 120-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/05/2025	
2	NDC:70108- 120-02	1 in 1 BOX	05/05/2025	
2		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.