SMART CARE SPIDER-MAN HAND SANITIZER- ethyl alcohol spray
Ashtel Studios, Inc.

Smart Care® MARVEL SPIDER-MAN 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, Blue 1, Denatonium Benzoate, Fragrance, Glycerin, Tocopheryl Acetate, Water, Yellow 5

Questions or comments?

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

KILLS 99.9% OF GERMS*

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

DESIGNED IN U.S.A.
MANUFACTURED IN CHINA

Smart Care®
WWW.SMARTCAREUS.COM

1-909-434-0911 INTERNATIONAL

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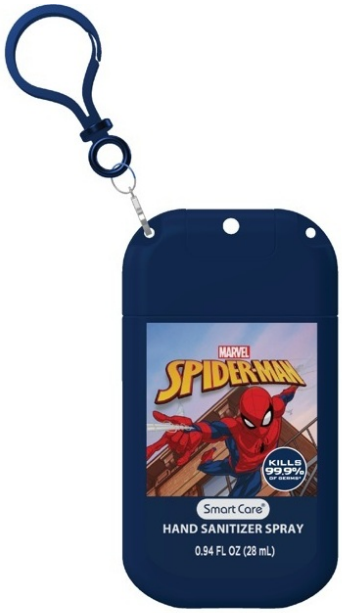
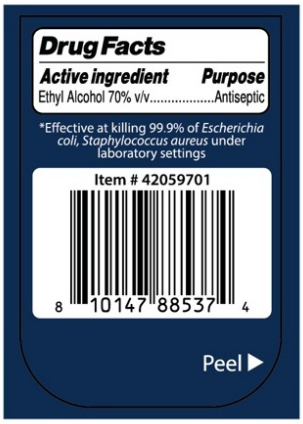
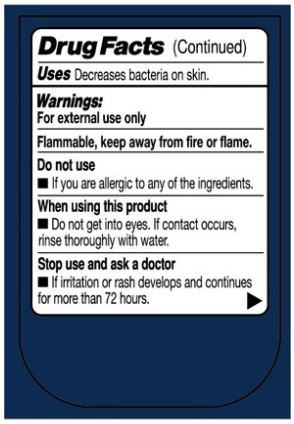
DISTRIBUTED BY ASHTEL STUDIOS INC.
ONTARIO, CALIFORNIA 91761

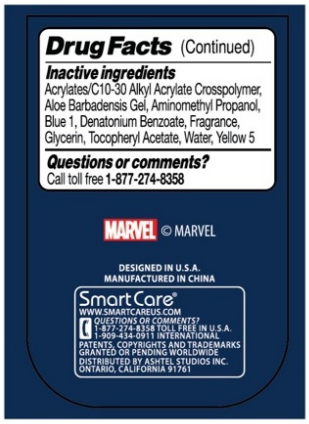
Packaging



Drug Facts Label







SMART CARE SPIDER-MAN HAND SANITIZER

ethyl alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70108-162
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)	
.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:70108-162-01	28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/28/2025	

Labeler - Ashtel Studios, Inc. (148689180)

Revised: 5/2025 Ashtel Studios, Inc.