

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

Flammability. 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 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

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

*Effective at killing 99.9% Escherichia coli, Staphylococcus aureus.

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**DESIGNED IN U.S.A.
MANUFACTURED IN CHINA**

Smart Care®

WWW.SMARTCAREUS.COM


1-909-434-0911 INTERNATIONAL

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ONTARIO, CALIFORNIA 91761

Packaging

\$5 SmartCare® **3PK**



MARVEL SPIDER-MAN

HAND SANITIZER SPRAY
THREE 0.94 FL OZ (28 mL) • TOTAL 2.82 FL OZ (84 mL)

*Effective at killing 99.9% Escherichia coli, Staphylococcus aureus.

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ONTARIO, CALIFORNIA 91761

Item # 88113
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SMART CARE SPIDER-MAN HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70108-086
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:70108-086-01	3 in 1 BOX	06/15/2024	
1		28 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)	06/15/2024	

Labeler - Ashtel Studios, Inc. (148689180)

Revised: 6/2024

Ashtel Studios, Inc.