SMART CARE HAND SANITIZER- ethyl alcohol gel
Ashtel Studios, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Smart Care® Hand SANITIZER

Drug Facts

Active Ingredients

Ethyl Alcohol 62% v/v

Purposes

Antiseptic

Use:

To help reduce bacteria and germs on the skin.

Warnings:

For external use only: hands

Flammable, keep away from fire or flame.

When using this product

• Keep out of eyes.

• In case of contact with eyes, flush thoroughly with water.

• Avoid contact with broken skin.

• Do not inhale or ingest.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions:

Put a thumbnail size amount in your palm and rub your hands together briskly until dry. Children under 6 years of age should be supervised when using this product. Not recommended for infants.

Other Information:

• Do not store above 100° F (38°C)

• May discolor some fabrics.

• Harmful to wood finishes and plastics.

Inactive Ingredients:

Water (Aqua), Glycerin, Propylene Glycol, Carbomer, Fragrance, Triethanolamine, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate (Vitamin E), FD&C Yellow No.5, FD&C Blue No.1.

KILLS UP TO 99.99% OF MOST COMMON GERMS

Aloe Vera & Vitamin E

FAST & EFFECTIVE

Keep bottle of Hand Sanitizer in:

- Automobile
- Bathrooms
- Workplace
- Kitchen
- Medical Offices
- Nurseries

SMARTCAREUS.COM

QUESTIONS OR COMMENTS?

1-877-274-8358 TOLL FREE IN U.S.A.

1-909-434-0911 INTERNATIONAL

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DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

DESIGNED IN U.S.A.

MADE IN CHINA

Packaging

SMART CARE HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70108-056

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70108-056-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020	
2	NDC:70108-056-02	236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020	
3	NDC:70108-056-03	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020	
4	NDC:70108-056-04	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/08/2020	

Labeler - Ashtel Studios, Inc (148689180)

Revised: 6/2020 Ashtel Studios, Inc