SMART CARE HAND SANITIZER 8 OZ- 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%

Purposes

Antiseptic

Uses

For external use only: hands

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 SMART CARE® HAND SANITIZER. 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), Aloe Barbadensis Leaf Extract, Carbomer, Parfum, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Triethanolamine, FD&C Yellow No. 5(Tartrazine), FD&C Blue No. 1

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.

KILLS UP TO 99.9% OF MOST COMMON GERMS

Aloe Vera & Vitamin E

FAST & EFFECTIVE

SMARTCAREUS.COM

QUESTIONS OR COMMENTS?

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

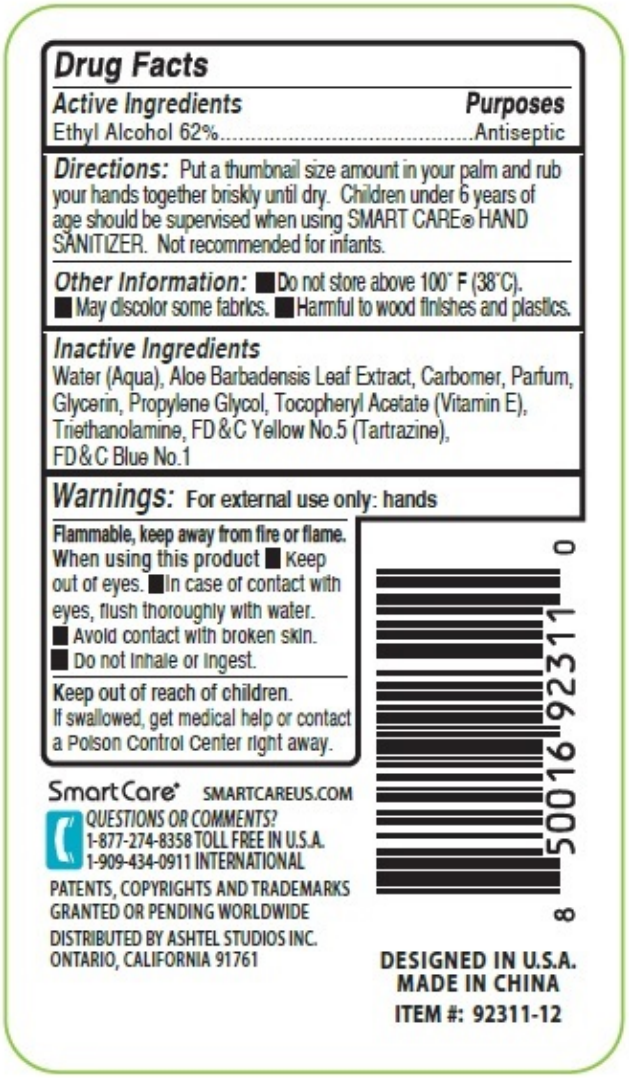
1-909-434-0911 INTERNATIONAL

PATENTS, COPYRIGHTS AND TRADEMARKS GRANTED OR PENDING WORLDWIDE

DISTRIBUTED BY ASHTEL STUDIOS INC. ONTARIO, CALIFORNIA 91761

DESIGNED IN U.S.A. • MADE IN CHINA

Packaging



SMART CARE HAND SANITIZER 8 OZ

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70108-049
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)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
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:70108-049-01	236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/06/2020	

Labeler - Ashtel Studios, Inc (148689180)

Revised: 4/2020 Ashtel Studios, Inc