ANTIBACTERIAL HAND SANITIZER- ethyl alcohol gel Hit Promotional Products, 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.

Antibacterial Hand Sanitizer

Active Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

For Hand Sanitizing

Warnings

For external use only - hands.

Flammable. Keep away from heat and 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 call a Poison Control Center right away.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.
- For children under 6 age use only under adult supervision.
- Not recommended for infants.

Other Information

- Do not store above 105F.
- May discolor some fabrics.

Inactive Ingredients

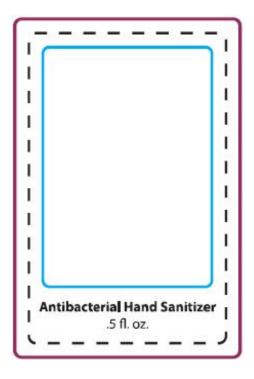
Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate, Triethanolamine, Water. Yellow 5 (CI 19140), Blue 1 (CI 42090)

Principal Display Panel

Antibacterial Hand Sanitizer

.5 fl. oz.

0.5oz #9067 Hit® ASI-61125 HIT Promotional Product Inc. Largo, FL. Made in China



Label

ANTIBACTERIAL HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70204-504
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBO XYPO LYMETHYLENE (UNII: 0 A5MM307FC)	

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TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70204-504-01	14.8 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/07/2020	
2 NDC:70204-504-02	29.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/07/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/07/2020	

Labeler - Hit Promotional Products, Inc. (081070202)

Revised: 5/2020 Hit Promotional Products, Inc.