

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), Yellow 6 (CI 15985)

Principal Display Panel

Antibacterial Hand Sanitizer

.5 fl. oz.

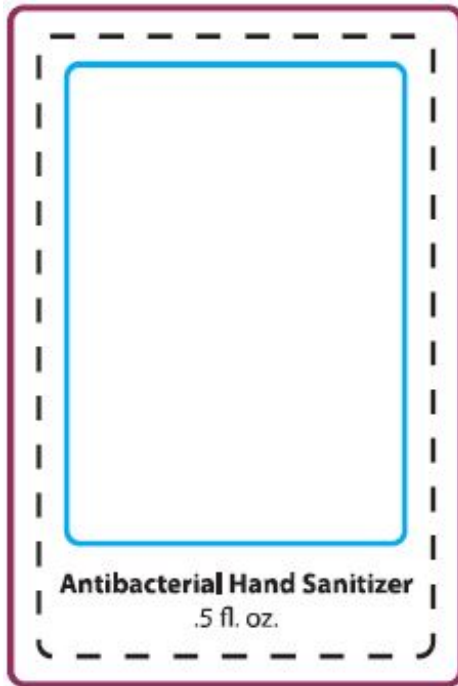
0.5oz #9067

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:70204-506
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)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	

TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70204-506-01	14.8 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/07/2020	
2	NDC:70204-506-02	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.