

**BODY PRESCRIPTIONS VANILLA ANTIBACTERIAL HAND SANITIZER- ethyl alcohol spray**  
**ENCHANTE ACCESSORIES 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.*

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**50563-417/JAS21AS4**

**Active Ingredient**

Alcohol Denat: 75%

**Purpose**

Antiseptic

**Use**

- To decrease bacteria on the skin that could cause disease
- Recommended for repeated use

**WARNINGS**

For external use only

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

**Stop use and ask a doctor** if skin irritation develops

**Keep out of reach of children.** In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately

**Directions**

Wet hands thoroughly with product and allow to dry without wiping

For children under 6, use only under adult supervision

Not recommended for infants

## Other information

Do not store above 105F

May discolor some fabrics

Harmful to wood finishes and plastics

## Inactive ingredients

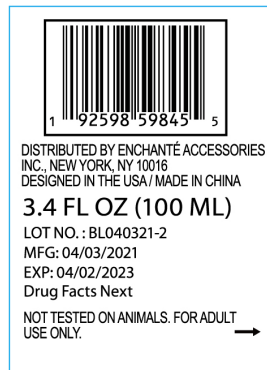
Water, Fragrance, Red 4, Yellow 5

10346-2-6-10 (JAS21AS4)

40\*60mm



40\*55mm



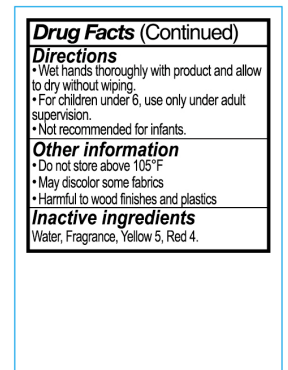
first page

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second page

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third page

## BODY PRESCRIPTIONS VANILLA ANTIBACTERIAL HAND SANITIZER

ethyl alcohol spray

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50563-417
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>WATER</b> (UNII: 059QF0KO0R)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	

<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50563-417-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2021	

<b>Marketing Information</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/25/2021	

**Labeler** - ENCHANTE ACCESSORIES INC. (186050696)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Xiamen Afgamarine Biotechnology Co., Ltd.		420485152	manufacture(50563-417)

Revised: 3/2021

ENCHANTE ACCESSORIES INC.