# BODY PRESCRIPTIONS VANILLA INFUSED WITH VITAMIN E ANTIBACTERIAL HANDbenzalkonium chloride lotion 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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#### 50563-335/JAS21F16

# **Active Ingredient**

Benzalkonium Chloride 0.13%

# Purpose

Antibacterial

## Use

■ To help reduce the bacteria on the skin

## **WARNINGS**

For external use only

When using this product, keep out of eyes. In case of contact with eyes, flush thoroughly with water.

Stop use and ask a doctor if skin irritation develops.

Keep out of reach of children. In case of accidenta ingestion, seek professional assistance or contact a Poison Control Center immdiately.

## Directions

Apply a small amount to palms and rub hands together thoroughly

## Other Information

Do not store above 105F

May doscolor some fabrics

Harmful to wood finishes and plastics

## **Inactive ingredients**

Water, Cetearyl Alcohol, Mineral Oil, Glyceryl Stearate, PEG 100 Stearate, Glycerin, C13-14 Isoparaffin, Laureth-7, Polyacrylamide, Phenoxyethanol, Benzyl Alcohol, Fragrance (Parfum), Butyrospermum Parkii (Shea) Butter, Citric Acid, Tocopheryl Acetate, Yellow 5, Red 4

#### 189\*70mm

#### **BODY PRESCRIPTIONS**

ANTI-BACTERIAL

# HAND LOTION

## Vanilla

INFUSED WITH VITAMIN E

kills bacteria soften · hydrates · protects · moisturizing

8 fl oz (236.6 ml)



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NOT TESTED ON ANIMALS. FOR ADULT USE ONLY.
DESIGNED IN THE USA / MADE IN CHINIA
FORMULATED EXCLUSIVELY AND DISTRIBUTED BY:
ENCHANTÉ ACCESSORIES INC., NEW YORK, NY 10016
8 FL OZ (236.6 ML)

LOT NO. : BL091020-5 MFG: 09/10/2020 EXP: 09/09/2022



# BODY PRESCRIPTIONS VANILLA INFUSED WITH VITAMIN E ANTIBACTERIAL HAND

benzalkonium chloride lotion

### **Product Information**

HUMAN OTC DRUG Product Type Item Code (Source) NDC:50563-335

**Route of Administration TOPICAL** 

I	Active Ingredient/Active	Moiety
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**Basis of Strength** Strength Ingredient Name BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -BENZALKONIUM 0.13 g UNII:7N6JUD5X6Y) **CHLORIDE** in 100 mL

**Inactive Ingredients Ingredient Name** Strength MINERAL OIL (UNII: T5L8T28FGP) GLYCERYL STEARATE SE (UNII: FCZ5MH785I) PEG-100 STEARATE (UNII: YD01N1999R) C13-14 ISOPARAFFIN (UNII: E4F12ROE70) GLYCERIN (UNII: PDC6A3C0OX) WATER (UNII: 059QF0KO0R) POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV) **LAURETH-7** (UNII: Z95S6G8201) PHENOXYETHANOL (UNII: HIE492ZZ3T) BENZYL ALCOHOL (UNII: LKG8494WBH) BENZOIC ACID (UNII: 8 SKN0 B0 MIM) CITRIC ACID MONOHYDRATE (UNII: 2968 PHW8 QP) BUTYRO SPERMUM PARKII (SHEA) BUTTER UNSAPO NIFIABLES (UNII: 0 C9 AC7D6 XU) .ALPHA.-TO COPHEROL ACETATE (UNII: 9E8X80D2L0) FD&C RED NO. 4 (UNII: X3W0 AM1JLX)

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1 NDC:50563-335-01 236.6 mL in 1 BOTTLE; Type 0: Not a Combination Product 10/11/2020

# **Marketing Information**

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/11/2020	

# Labeler - ENCHANTE ACCESSORIES INC. (186050696)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
Xiamen Afgamarine Biotechnology Co., Ltd.		420485152	manufacture (50563-335)			

Revised: 10/2020 ENCHANTE ACCESSORIES INC.