## SANITELLE MOISTURIZING HAND SANITIZER- alcohol gel BENTUS LABORATORII, OOO

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.

-----

#### **Active Ingredient(s)**

Alcohol 80% v/v. Purpose: Antiseptic

#### **Purpose**

Antiseptic, Hand Sanitizer

#### Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

#### Warnings

For external use only. Flammable. Keep away from heat or flame

#### Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

#### Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

#### **Inactive ingredients**

Water, Propylene Glycol, Glycerin, , Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine, Tocopherol

#### Package Label - Principal Display Panel







50 mL NDC: 80068-005-50







250 mL NDC: 80068-005-25

Distibuted by: Flex Technologies, Inc, 10432 Balls Ford Rd. Suite 300, Manassas, VA

20109, USA

# SANITELLE MOISTURIZING HAND SANITIZER alcohol gel

## Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:80068-005

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	66.2 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
TOCOPHEROL (UNII: ROZB2556P8)	
GLYCERIN (UNII: PDC6A3C0OX)	0.79 mL in 100 mL
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 903K93S3TK)	

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:80068- 005-50	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/23/2021		
2	NDC:80068- 005-25	250 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/23/2021		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/19/2021	

### Labeler - BENTUS LABORATORII, OOO (354757383)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
BENTUS LABORATORII, OOO		354757383	manufacture(80068-005)	

Revised: 2/2021 BENTUS LABORATORII, 000