# LAVENDER- hand sanitizer liquid Sante Manufacturing 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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Active Ingredients - Ethyl Alcohol 70%

Purpose - Antiseptic

#### Use

Use hand sanitizer to help reduce disease causing bacteria on the skin

Warnings

Flammable. Keep away from fire or flame

For external use only -

**Avoid contact with eyes**. In case of contact, rinse eyes thoroughly with water

**Stop use and ask a doctor if** irritation or redness develops

#### Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- apply liberally to hands and rub thoroughly until dry
- supervise children in teh use of this product

Water, Carbomer, Triethanolamine, Glycerin, Propylene Glycol, Lavandula Angustifolia Flower / Leaf / Steam Extract, Fragrance, D&C Red#33, FD&C Blue # 1

SpaSoap

Hand Sanitizer

Lavender

20z / 59ml





#### LAVENDER

hand sanitizer liquid

| Droduct  | Information |
|----------|-------------|
| Praniiri | Intormation |

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71020-012

Route of Administration TOPICAL

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

Ingredient Name Basis of Strength Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)

ALCOHOL (UNII: 3K9958V90M) ALCOHOL 700 mL in 1000 mL

### **Inactive Ingredients**

| Ingredient Name                                     | Strength |  |
|---|----------|--|
| CARBOMER 940 (UNII: 4Q93RCW27E)                     |          |  |
| GLYCERIN (UNII: PDC6A3C0OX)                         |          |  |
| LAVANDULA ANGUSTIFO LIA FLO WER (UNII: 19 AH1RAF4M) |          |  |
| TRIETHANOLAMINE LAURYL SULFATE (UNII: E8458C1KAA)   |          |  |

| WATER (UNII: 059QF0KO0R)           |  |
|------------------------------------|--|
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD) |  |
| D&C RED NO. 33 (UNII: 9DBA0SBB0L)  |  |

| l | Packaging              |   |                         |                       |
|---|------------------------|---|-------------------------|-----------------------|
|   | # Item Code            | Package Description   | Marketing Start<br>Date | Marketing End<br>Date |
|   | 1 NDC:71020-012-<br>02 | 59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product | 03/11/2017              |                       |

| Marketing Information   |  |                      |                    |  |
|-------------------------|--|----------------------|--------------------|--|
| Marketing Category      | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |  |
| OTC monograph not final | part333A                                 | 03/11/2017           |                    |  |
|                         |  |                      |                    |  |

## Labeler - Sante Manufacturing Inc (242048747)

| Establishment           |         |           |                        |  |  |
|-------------------------|---------|-----------|------------------------|--|--|
| Name                    | Address | ID/FEI    | Business Operations    |  |  |
| Sante Manufacturing Inc |         | 242048747 | manufacture(71020-012) |  |  |

Revised: 3/2017 Sante Manufacturing Inc