FIRST HAND HAND SANITIZER- alcohol liquid South Tenth Development, LLC

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.

"First Hand" Hand Sanitizer

DRUG FACTS

Active Ingredient

Ethyl Alcohol 80%

Purpose

Antiseptic

Uses

Reduce bacteria on the skin. Recommended for repeated use

Warnings

For external use only.

Flammable Keep away from fire, flame, or sparks

When using this product

do not use in or near eyes. In case of contact, rinse eyes thoroughly with water

Stop use & ask a doctor

If irritation or rash appears on the skin.

Keep out of reach of children.

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

Directions

- Place enough to cover both hands in the palm, and rub hands together until dry.
- Children under 6 years of age should be supervised by an adult when applying this product.

Other Information

- Store below 105°F.
- May discolor certain fabrics or surfaces.

Inactive Ingredients

Distilled water, glycerol, hydrogen peroxide, denatonium benzoate.

Package Labeling

Trim Bleed Print-to-Die Tolerance



FIRST HAND HAND SANITIZER

alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:78705-000

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)ALCOHOL0.8 mL in 1 mL

Inactive Ingredients

| indente ingredients | | | | |
|--|----------|--|--|--|
| Ingredient Name | Strength | | | |
| WATER (UNII: 059QF0KO0R) | | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | | |
| HYDRO GEN PERO XIDE (UNII: BBX060AN9 V) | | | | |
| DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2) | | | | |

Packaging

| | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|--|---|-----------|---------------------|-----------------------------|--------------------|
|--|---|-----------|---------------------|-----------------------------|--------------------|

1 NDC:78705-000-00 295 mL in 1 BOTTLE; Type 0: Not a Combination Product 06/01/2020

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|-------------------------|--|----------------------|--------------------|
| OTC monograph not final | part333E | 06/01/2020 | |
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Labeler - South Tenth Development, LLC (117521920)

Revised: 6/2020

South Tenth Development, LLC