ANTIBACTERIAL HAND SANITIZER- alcohol spray National Design 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.

Antibacterial hand sanitizer

Drug Facts

Active ingredient

Ethyl Alcohol 68%

Purpose

Antiseptic

Uses

• For hand sanitizing

Warnings

For external use only hands.

Flammable. Keep away from heat or flame.

When using this product

Keep out of eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin.

Keep out of reach of children

Other information:

store at 20°C (68° to 77° F). may discolor fabrics.

Ingreidients:

Glycerin, propylene glycol, triethanolamine, carbomer, fragrance, disodium edta, tocopheryl acetate, deionized water.

Package Labeling:

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deionized water.

Factoy Information:

Ningbo Lanzi Cosmetic Technology Co.,Ltd

MADE IN CHINA



DRAWINGS SCHOOL

Antibacterial Hand Sanitizer 0.36 oz. / 10 ml.

ANTIBACTERIAL HAND SANITIZER

alcohol spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:80575-000

Route of Administration TOPICAL

Active Ingredient/Active Moiety

| ingredient Name | basis of Strength | Strength |
|---------------------------------------------------------|-------------------|-----------------|
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) | ALCOHOL | 0.68 mL in 1 mL |

Inactive Ingredients

| ı | Ingredient Name | Strength |
|---|-------------------------------------|----------|
| ı | GLYCERIN (UNII: PDC6A3C0OX) | |
| ı | PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | |
| | | |

TROLAMINE (UNII: 9O3K93S3TK)

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)

| EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM) | |
|----------------------------------------------------|--|
| .ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0) | |
| WATER (UNII: 059QF0KO0R) | |

| Packaging | | | |
|----------------------|-------------------------------------------------------------|-------------------------|-----------------------|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| NDC:80575-000- 01 | 10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 09/01/2020 | |

| Marketing Information | | | |
|-------------------------|------------------------------------------|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | part333E | 09/01/2020 | |
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Labeler - National Design LLC. (360763320)

Revised: 9/2020 National Design LLC.