HAND SANITIZER CLEAR- alcohol gel Panros a Enterprises, 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.

Hand Sanitizer Clear

Drug Facts

Active ingredients

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- hand sanitizer to help decrease bacteria on the skin
- when water, soap & towel are not available
- recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

Do not apply around eyes.

Do not use

in ears & mouth

When using this product,

avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor if

redness or irritation develop and persist for more than 72 hours.

Keep out of reach of children.

Children must be supervised in use of this product.

Directions

- pump as need into your palms and thoroughly spread on both hands.
- rub into skin until dry.

Other information

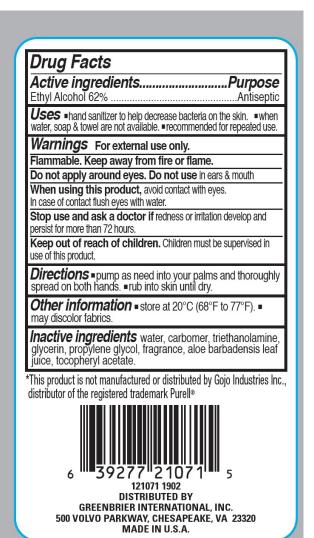
- store at 200 (680 to 770).
- may discolor fabrics.

Inactive ingredients

water, carbomer, triethanolamine, glycerin, propylene glycol, fragrance, aloe barbadensis leaf juice, tocopheryl acetate.

Package Labeling:





HAND SANITIZER CLEAR

alcohol gel

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:50302-000 |
| Route of Administration | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|--|-------------------|----------------|
| Ingredient Name | Basis of Strength | Strength |
| ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) | ALCOHOL | 620 mg in 1 mL |
| | | |

| Inactive Ingredients | | | |
|--|----------|--|--|
| Ingredient Name | Strength | | |
| WATER (UNII: 059QF0KO0R) | | | |
| CARBO XYPO LYMETHYLENE (UNII: 0 A5MM307FC) | | | |
| TROLAMINE (UNII: 9O3K93S3TK) | | | |
| GLYCERIN (UNII: PDC6A3C0OX) | | | |
| PROPYLENE GLYCOL (UNII: 6DC9Q167V3) | | | |
| ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X) | | | |

| l | Packaging | | | | |
|---|-----------|-----------------|---|-----------------------------|--------------------|
| l | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| l | 1 NI | DC:50302-000-00 | 237 mL in 1 BOTTLE; Type 0: Not a Combination Product | 03/01/2019 | |

| Marketing Information | | | |
|-------------------------|--|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC monograph not final | part333A | 0 3/0 1/20 19 | |
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Labeler - Panrosa Enterprises, Inc. (859957578)

Revised: 2/2019 Panrosa Enterprises, Inc.